The immediate early protein 1 of the human herpesvirus 6B counteracts NBS1 and prevents homologous recombination repair pathways

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Keywords: DNA double-strand break signaling, telomere; integration; human herpesvirus 6A/B; immediate-early protein IE1.

Abstract

Integration of viral DNA in the genome of host cells triggers host-pathogens interaction that are consequential for the virus and the infected cells. In cells semi-permissive for viral replication, the human herpesvirus 6B (HHV-6B) integrates its genome into the host telomeric sequences. Interestingly, HHV-6B integration in gametes leads to a condition called inherited chromosomally integrated HHV-6B (iciHHV-6B), where the newborn carries a copy of HHV-6B in every cell of its body and is associated with health issues such as spontaneous abortion rates, pre-eclampsia and angina pectoris when transmitted to its offspring. Unlike retroviruses, the mechanism that leads to viral integration of DNA viruses and the consequences of these events on host cells are not well characterized. Here, we report that HHV-6B infection induce genomic instability by suppressing the ability of the host cell to sense DNA double-strand break (DSB). We discovered that this phenotype is mediated by the ability of the immediate-early HHV-6B protein IE1 to bind, delocalize, and inhibit the functions of the DNA damage sensor NBS1. These results raise the possibility that the genomic instability induced by the expression of IE1 from integrated genomes contributes to the development of iciHHV-6B-associated disease. As reported for other types of viruses, the inhibition of DSB sensing and signaling promotes viral replication. However, HHV-6B integration is not affected when this pathway is inhibited, supporting models where integration of the viral genome at telomeric sequence is dictated by mechanisms that promote telomere-elongation in a given infected cell and not solely DNA repair mechanisms.

Introduction

Human herpesvirus 6B (HHV-6B) is a betaherpesvirus that infects nearly 90% of the world's population in the first two years of life and is responsible for Roseola Infantum, a pathology defined by skin rashes, high fevers and respiratory distress(1–3). In this DNA virus subfamily, HHV-6B shares 94% homology with HHV-6A, another lymphotropic virus. Like other herpesviruses, HHV-6A and HHV-6B (HHV-6A/B) establish lifelong latency in infected hosts and can reactivate occasionally(4). However, whereas most herpesviruses achieve latency through the circularization and silencing of their genome, HHV-6A/B viruses can integrate their genome in the host chromosome terminal repeats called telomeres (chromosomally integrated HHV-6B (ciHHV-6B))(5, 6). If HHV-6A/B integration occurred in gametes before fertilization, the newborn carries a copy of HHV-6A/B in every cell of its body and can transmit it to its offspring. This condition called inherited chromosomally integrated HHV-6A/B (iciHHV-6A/B) concerns ~1% of the world's population, representing almost 80 million people(7, 8). iciHHV-6A/B is more prevalent in individual suffering from health issues such as spontaneous abortion rates(9), pre-eclampsia(10) and angina pectoris(11) compared to healthy subjects (reviewed in(12, 13)). However, consequences of iciHHV-6A/B are still poorly understood due to lack of clinical associations.

The linear double-strand DNA (dsDNA) genome of HHV-6A/B is flanked by an array of direct repeats containing 15 to 180 reiterations of 5'-TTAGGG-3' perfect telomeric repeats (pTMRs) that are identical to human telomeric sequences, and which are important for viral integration(14). HHV-6A/B genome integration occurs at telomeres in a process that is dependent on the integrity of these pTMRs(14). Based on this observation, current models propose that viral integration is mediated through homology-directed repair (HDR) processes including single stranded annealing (SSA) or break-induced replication (BIR)(15). These HDR pathways are favored given that the integration occurs in an oriented manner that is driven by one of the pTMR(14, 16). In this case, HHV-6A/B genome would integrate at sites of DNA double-stranded break (DSB) that are caused following replication fork collapse at telomeres upon replication stress. In both scenarios, the annealing of the pTMR sequence to the 3' overhang generated by partial resection of DNA ends of the telomere would drive integration(15, 17).

In mammalian cells, HDR uses homologous sequences as template to repair breaks in a faithful manner. During this process, broken DNA ends are first detected by the MRN complex (MRE11, RAD50, NBS1)(18). The accumulation of the MRN complex at the break induces a signaling cascade that leads to the activation of the serine–threonine kinase ataxia–telangiectasia mutated (ATM) and concomitant phosphorylation of the histone variant H2AX on Ser139 (γ -H2AX). The interaction of MDC1 with γ -H2AX then triggers the ubiquitylation of the chromatin that surrounds the break by promoting the accumulation of the E3- ubiquitin ligases RNF8 and RNF168(19, 20). In S/G2 phase of the cell cycle, the recruitment of the DNA repair factor BRCA1 and the nuclease CtIP to ubiquitylated chromatin cooperates with EXO1 and BLM-DNA2 nucleases to facilitate extensive end resection. Extensive accumulation of single-strand DNA (ssDNA) is generated through this process or by uncoordinated DNA unwinding, and DNA synthesis that occurs at stalled replication forks ultimately leading to the recruitment of recombinases that drive homology search(21). RAD51 or RAD52 recombinases promote DNA repair by HDR and SSA(22). Both recombinases also promote DNA repair of one-end DSB, but their exact contribution to that latter pathway is still unclear(23).

HHV-6A and B are two distinct viruses that share 90% sequence homology. Although they both have a tropism for CD4⁺ T lymphocytes, they present epidemiological, biological, and immunological differences(17). HHV-6B, which infects nearly 90% of the population in the first two years of life, is much more characterized than HHV-6A. HHV-6B expresses sequentially more than 97 proteins during its lytic cycle. Immediate early (IE) proteins are expressed early in the viral cycle and exhibit functions that regulate viral genes expression and promote the establishment of a favorable environment for infection. Interestingly, immediate early protein 1 (IE1) transcripts are detected in RNA-seq analysis on tissues extracted from iciHHV-6B+ individuals(24), suggesting that IE1 is expressed during latency. IE1 is the first protein expressed following HHV-6B infection(25) and although it is known to control the antiviral immune response by compromising type I interferon production and signaling(26, 27), its role during infection and the integration of the viral genome is still poorly defined. In infected cells, IE1 is exclusively localized within promyelocytic leukemia (PML) nuclear bodies (PML-NBs) (28), a nuclear structure that was recently implicated in DNA repair mediated by HDR through a yet undefined mechanism(29–32). Interestingly, depletion of PML reduces HHV-6B integration(33), suggesting that the IE1/PML-NBs may participate to viral integration.

In this study, we found that viral infection, and more specifically the expression of IE1, leads to the accumulation of micronuclei and numerous DSBs in cells. Further investigations revealed that the viral protein specifically prevents H2AX phosphorylation through a bipartite mechanism that relies on the ability of IE1 to interact with NBS1 and inhibit its interaction with ATM. While this function is independent of PML, structure function analysis identified a NBS1-interacting domain (NID) as well as NBS1-inhibitory domain (NBS1i) in the N-terminus and the C-terminus regions of IE1, respectively. Although current models propose that viral integration occurs through HDR DNA repair, we show that the expression of IE1 strongly inhibits all types of repairs that rely on homology. We show that both viral replication and integration are not affected by the depletion of NBS1 in cells where telomeres are elongated in a human telomerase reverse transcriptase (hTERT) dependent manner, a finding that is consistent with a role of the telomerase complex in this process(34). In contrast, in cells that rely on alternative telomere lengthening (ALT) mechanisms involving HDR events, knockdown of NBS1 negatively affected HHV-6B integration. Thus, in addition to identifying a new bipartite mechanism for the inhibition of NBS1 by a viral protein, our findings reveal that viral integration relies on biological pathways that safeguard telomere extension in infected cells. Importantly, as IE1 expression has been detected in cells where HHV-6A/B is integrated(35), our results suggest a potential role of genomic instability in the development of diseases associated with iciHHV-6A/B.

Results

HHV-6B infection and IE1 expression induce genomic instability

Infection of the lymphoblast T cell line MOLT-3 (permissive for viral replication) by HHV-6B rapidly induces the formation of micronuclei (MNi), suggesting that the virus leads to genomic instability during infection (Fig.1A and *SI Appendix*, Fig.S1A). Among the early HHV-6B proteins that are expressed upon infection, expression of IE1 in U2OS cell line is sufficient to promote the accumulation of MNi over time (Fig. 1B and *SI Appendix*, Fig.S1B-C). Such MNi are compartmentally separated from the primary nucleus that are surrounded

by an envelope (*SI Appendix*, Fig. S1D-E). They arise from unresolved genomic instability such as DSBs (i), lagging chromosome (ii) or by the rupture of anaphase bridges (ABs) (iii) (Fig. 1C)(36). Further analysis of the IE1-induced MNi revealed that a much lower proportion of these MNi exhibit centromere staining (Fig. 1D and *SI Appendix*, Fig. S1F), suggesting that they are not induced by chromosome segregation defects. Although IE1 colocalizes with telomeres(37), fluorescence *in situ* hybridization revealed that IE1-induced MNi accumulate similar levels of telomeric DNA than the micronuclei observed in parental U2OS cell (Fig. 1E and *SI Appendix*, Fig. S1G). Moreover, IE1 is only detected in approximately 5-10% of these micronuclei (*SI Appendix*, Fig. S1H), suggesting that micronuclei are not arising from IE1-induced genomic instability at telomeres. Interestingly, metaphase spread assays revealed that IE1-expressing cells exhibit higher frequency of DNA breaks (Fig. 1F-G), supporting the hypothesis that the micronuclei accumulation results from the accumulation of DSBs.

HHV-6B impairs DSB-signaling by interacting with NBS1

Accumulation of DSBs is either caused by increased source of DNA breaks or by defective DNA DSB-signaling and repair. To determine how IE1 promotes genomic instability, we first investigated whether U2OS clones that stably express the viral protein accumulate the DSB γ -H2AX marker. Surprisingly, γ -H2AX is dramatically reduced following exposure to irradiation (IR) in cells that express IE1 (Fig. 2A-B). This inhibition is independent of the accumulation of IE1 within PML-NBs as a similar phenotype is observed in PML-deficient U2OS that transiently express IE1 (PML^{-/-}, Fig. 2C and *SI Appendix*, Fig. S2A-C). Importantly, the inhibition of γ -H2AX by IE1 is recapitulated in irradiated HHV-6B-infected MOLT-3 cells (Fig. 2D-E), indicating that DSB-signaling is also impaired by IE1 in the context of a natural infection.

The histone variant H2AX is phosphorylated at DSBs following the activation of ATM by the MRN complex(18). To determine how IE1 interferes with DSB-signaling, we first investigated the localization of NBS1 and MRE11 in U2OS clones expressing IE1. In the absence of irradiation, both NBS1 and MRE11 proteins colocalize with IE1 foci (Fig. 3A-D and *SI Appendix*, Fig. S3A). Interestingly, only NBS1 is relocated to IE1 foci following irradiation suggesting that the recruitment of MRE11 in absence of irradiation is mediated by NBS1 (*SI Appendix*, Fig. S3B-C). Consistent with this model, the recruitment of MRE11 to IE1 foci is impaired in NBS1-depleted U2OS cells that transiently express IE1 (*SI Appendix*, Fig. S3D-F). Furthermore, when mCherry-LacRnls-IE1 fusion protein is recruited to a *LacO* array in U2OS 2-6-5 transfected cells(38, 39), only NBS1 is recruited to the array with an efficiency similar to its recruitment to DSBs induced by the ER-mCherry-LacR-FOKI-DD endonuclease, which is used as a positive control in this assay (Fig. 3E-F and *SI Appendix*, Fig. S3G-H). The absence of γ -H2AX signaling at the array upon the recruitment of mCherry-LacRnls-IE1 shows that the viral protein recruits NBS1 independently of DSB signaling. As observed in Fig. 2, the interaction of IE1 with NBS1 is independent of its localization to PML-NBs as NBS1 is recruited to IE1 foci with a similar efficiency in PML^{-/-} U2OS cells (*SI Appendix*, Fig. S3I-J). Altogether, these results suggest that IE1 recruits NBS1 and inhibits γ -H2AX signaling by preventing its recruitment to endogenous DNA breaks.

Identification of NBS1 bipartite interaction and inhibition domain in IE1

The functional domains of IE1 are not well characterized aside from a STAT2 binding domain that was mapped in the N-terminal domain of the protein (amino acids 270-540)(26). Guided by a secondary structure analysis of the protein, a series of IE1 fragments were fused to mCherry-LacRnls to assess their ability to recruit endogenous NBS1 to the *LacO* array (Fig. 4A). Using this approach, we observed that the fragment comprising amino acids (aa) 1-540 is sufficient to recapitulate the level of NBS1 recruitment observed with the full-length protein (Fig. 4B and *SI Appendix*, Fig. S4A-B). In this assay, the C-terminal domain of the viral protein aa 541-1078 was also able to partially recruit NBS1 to the *LacO*-array. As the fragment composed of aa 541-809 does not promote NBS1 recruitment, we concluded that the domain 810-1078 also interacts with NBS1. Interestingly, we found that only WT and the C-terminal (aa 541-1078) domain of IE1 were able to inhibit γ -H2AX signaling in irradiated U2OS cells (Fig. 4C-D and *SI Appendix*, Fig. S4C). Together, these results suggest that IE1 interacts and inhibits NBS1 using bipartite motifs. The N-terminal of IE1 is composed of a NBS1-interacting domain (NID) and the C-terminal domain independently inhibits the ability of NBS1 to activate ATM (Fig. 4A). Based on this observation, we named this C-terminal domain: NBS1 inhibitory domain (NBS1i).

NBS1 encodes a 95 kDa protein that contains multiple domains that are required for its recruitment to DSBs and its interaction with the PI3K ATM and ATR(40). Briefly, NBS1 contains a forkhead-associated (FHA) and two breast cancer C-terminal domains (BRCTs) that are both required for optimal phospho-dependent accumulation of the protein at the break. The C-terminal part of the protein contains a domain that promotes its interaction with MRE11 (MRE11-binding motif, MBM) and ATM (ATM-binding motif, ABM) (Fig. 4F). In the *LacO*-LacR assay, the recruitment of mCherry-LacRnls NBS1 to the array is sufficient to promote the phosphorylation of γ -H2AX(41), a function that is dependent of its ability to interact with ATM (Fig. 4E). Consistent with the inhibitory function of IE1, expression of the viral protein is sufficient to inhibit NBS1-induced γ -H2AX signaling at the array (Fig. 4E and *SI Appendix*, Fig. S4F). In this system, the mCherry-LacRnls-NBS1 328-754 was unable to efficiently recruit IE1 to the array (Fig. 4F-G and *SI Appendix*, Fig. S4G), suggesting the IE1 interacts with the BRCT2 domain of NBS1. Furthermore, LacR constructs that only lack the linker region also exhibits reduced recruitment of IE1 to the array, suggesting that the interaction of IE1 with NBS1 also relies on the integrity of this regions. Thus, our results support a model where IE1 need to contact two regions, the BRCA2 domain and the linker region. Whether both the NID and the NBS1i contact these regions or whether this is only mediated by the NID is unknown.

IE1 inhibits HDR repair pathways

DSBs signaling is essential to trigger the activation of DNA repair pathways that have been proposed to drive HHV-6B integration(15). As the functions of NBS1 are essential to promote the resection of DNA ends that trigger homology-directed repair, we investigated whether IE1 interferes specifically with this process using a panel of well-characterized DNA repair reporter assays. The efficiency of pathways that rely on homology-based DNA repair was assessed using DR-GFP and CRISPR-LMNA assays (homologous recombination)(42, 43), a SA-GFP assay (single-strand annealing)(42) and a RAD51-dependent BIR-GFP assay (break-induced replication)(44) (Fig. 5A-C and *SI Appendix*, Fig. S5A, top panels). In all assays, a condition without the endonuclease I-SceI was used as a negative control and the percentage of fluorescent-positive cells obtain with I-SceI was set to 1. Consistent with the ability of IE1 to inhibit the function of NBS1, both transient and stable

expression of the viral protein drastically abolished all types of homology-directed DNA repair (Fig. 5A-C, lower panel, and *SI Appendix*, Fig. S5A-C). As the clonal BIR-GFP U2OS cell line was generated in this study using previously described BIR-GFP reporter plasmid(44), we used siRNAs against RAD51 and RAD52 as additional controls (*SI Appendix*, Fig. S5D-F)(23). As expected, BIR-GFP signal was specifically inhibited in cells depleted for RAD51(44). In contrast to homology-based DNA repair, IE1 only slightly decreased or increased DNA repair in reporter assays that assess the efficiency of NHEJ (NHEJ-GFP EJ7 Fig. 5D, and NHEJ-pc222 *SI Appendix*, Fig. S5G). Altogether, these results are consistent with a model where homology-based DNA repair is specifically inhibited in cells that express HHV-6B IE1 and raise the point that either integration occurs in a context when the expression of the viral protein is minimal, absent, or driven through a homology-independent mechanism.

Integration of HHV-6B relies on the pathway that safeguards telomere elongation

The MRN complex is commonly targeted by viruses to promote viral replication(45) however, the requirement of this complex for viral integration is unclear. HHV-6B infection leads to different outcomes depending on the nature of the infected cells (Fig. 6A). In permissive cells, the expression of viral proteins promotes viral replication (lytic state). In contrast, in semi-permissive cells, integration of the viral genome at host's chromosome telomere is favored. The factors that lead to the reactivation of the integrated viral genome are still misunderstood. The fact that HHV-6B IE1 evolved to inhibit the function of NBS1 raises the possibility that the DNA repair protein negatively impact viral replication and/or integration. To investigate the role of NBS1 in these processes, permissive cells (MOLT-3) and semi-permissive cells (U2OS, HeLa, and GM847) depleted or not for NBS1 were infected with HHV-6B (*SI Appendix*, Fig. S6A-D). In MOLT-3, increased replication is measured in cells depleted for NBS1, suggesting that the protein is detrimental for HHV-6B replication (Fig. 6B). Interestingly, our results suggest that viral integration in semi-permissive cells relies on the molecular mechanisms that drive telomere elongation in these cells. Indeed, integration was not affected by the depletion of NBS1 in cells where telomeres lengthening is secured by hTERT (Table 1). However, integration is significantly reduced upon NBS1-depletion in both cell lines that rely on Alternative Lengthening of Telomeres to maintain telomere length (Table 1). In ALT-positive (ALT+) cells, telomere maintenance occurs on break-induced telomere synthesis, a RAD51-independent homology-directed DNA repair pathway(46). Thus, our data support a model in which the mechanism of viral integration is dictated by the telomere lengthening pathway of the infected semi-permissive cell rather than a common mechanism only driven by viral protein. Importantly, our data also imply that the expression of IE1 must be repressed to promote integration in ALT+ cells.

Table 1. Importance of NBS1 for HHV-6B chromosomal integration in ALT + and – cells.

Cell lines	ALT status	shRNA	% of cells integrated HHV-6B ^a (n) ^b	P value ^c
HeLa	Negative	CTRL	0.96 (36320)	<2.2e ⁻¹⁶
		NBS1	6.11 (33280)	
GM847	Positive	CTRL	0.65 (21820)	<2.2e ⁻¹⁶
		NBS1	0.01 (18320)	
U2OS	Positive	CTRL	1.60 (20000)	<2.2e ⁻¹⁶

		NBS1	0.69 (21520)	
U2OS PML ^{-/-}	Positive	CTRL	0.71 (28220)	ns
		NBS1	0.78 (30460)	

^a mean of three independent cultures

^b total number of cells analyzed

^c Pearson's Chi-squared test with Yates' continuity correction

Discussion

In this study, we set out to understand two phenomena in HHV-6B; 1) the molecular mechanism by which HHV-6B induces genomic instability in infected cells, and 2) the role of HDR-mediated repair pathways in the establishment of the latent state in semi-permissive cells. Using a series of microscopy- and cytometry-based approaches to track the source of DNA breaks in infected cells and in cells expressing the immediate-early protein IE1, we discovered that in both conditions, DNA double-strand break signaling, and repair are strongly inhibited through interference with the recruitment of the DNA repair protein NBS1 at the breaks. Specifically, we defined the molecular mechanism by which IE1 triggers the redistribution of the MRN complex to IE1-PML NBs by using a single-cell assay in which the colocalization of DNA repair factors with mCherry-tagged viral proteins is restricted to an integrated *LacO* array. These findings revealed that IE1 specifically interacts with NBS1 through a NBS1-interacting domain (NID) that is located in the N-terminal part of the viral protein. Furthermore, our finding revealed that the activation of ATM by NBS1 is strongly inhibited by a NBS1-inhibitory domain (NBS1i) located in the C-terminal region of IE1, supporting a model where IE1 impairs the function of NBS1 through bipartite motifs. Consistently, we found that the expression of IE1 specifically abolishes NBS1-dependent DNA repair pathways by using an array of well-established DNA repair reporter assays. Altogether, our work argues against a model where viral integration is promoted solely by homology-based repair, but rather supports models where integration of the viral genome at telomeres is dictated by mechanisms that promote telomere-elongation in a given infected cell. Our work suggests that IE1 expression must be tightly regulated to enable viral integration in cells where telomeres are elongated by ALT. Finally, our results raise the possibility that expression of IE1 from integrated genomes might contribute to the development of iciHHV-6B associated disease by inducing genomic instability in these cells. Using an RNA-seq approach, Peddu et al. reported that the *IE1* gene is among the most abundantly expressed genes in a variety of tissues from iciHHV-6+ individuals(47). Spontaneous and inducible IE1 protein expression from integrated HHV-6 genomes was also documented(35). At present, diseases associated with iciHHV-6A/B status include increased spontaneous abortion rates(9), pre-eclampsia(10) and angina pectoris(11). Further characterization of the proteins expressed from integrated genomes as well as the disease associated with these conditions will be required to strengthen our understanding of the consequences associated with viral latency in iciHHV-6B subjects.

The functions of the MRN complex are required at DSBs, stalled replication forks, chromosome segregation and dysfunctional telomeres to safeguard genomic stability in cells(18, 48). Viruses have evolved different strategies to adapt to their host cell environment. Many of them developed specific mechanisms to manipulate DNA damage signaling to either promote viral processes such as replication and integration, or to protect the integrity of their genome upon infection(45). A classic example comes from adenoviruses, where the

MRN complex is targeted by multiple viral protein to inhibit its function. Specifically, E4-ORF3-dependent relocalization of MRN proteins and E4-ORF6/E1B-55K-dependent degradation of MRN components is essential to enable transduction and efficient viral replication(49–54). In contrast, other viruses such as adeno-associated virus (AAV) rely on the activity of the complex for the integration of their genomes at the human AAVS1 site(55) or to promote a DNA damage response that enhances infection levels (Herpes Simplex Virus 1 (HSV-1) and Human Papillomavirus (HPV))(56–59). While these findings demonstrate that some viruses hijack the function of DNA repair protein to support different steps of the infection, the mechanisms by which these processes benefit viral replication remain a long-standing mystery in the field(45). In this study, we report that depletion of NBS1 results in increased HHV-6B replication in permissive cells, suggesting that HHV-6B also evolved to interfere with NBS1, or with the MRN complex, to prevent undesired recognition of viral DNA as broken DNA. This makes sense considering that during viral replication, numerous double-stranded linear genomes, which can be perceived as broken DNA, are generated. Interestingly, our structure function analyses revealed that the interaction between IE1 and NBS1 is mainly driven by the BRCT2 domain (aa 201-326) of NBS1 and, to a lesser extent, by the linker region (aa 327-638). These findings thus revealed that IE1 interacts with a domain of NBS1 that is essential for its MDC1-dependent chromatin retention of NBS1 to DSB (BRCT2)(60–64), providing a rational for its ability to compete with the recruitment and function of the DNA sensor protein. Interestingly, it has been proposed that the viral protein HSV-1 ICP0 also interacts with NBS1 through a region that span the linker (aa 590-710) to redirect HDR to specific loci during the infection(65). Here, the ability of IE1 to actively inhibit the NBS1-dependent activation of ATM at the *LacO* demonstrates that this is not the case for IE1 unless the NBS1i domain is post-translationally regulated during the infection. In contrast to AAV integration at the AAVS1 locus, the integration of HHV-6B at telomere is not strictly dependent on the MRN complex but rather on processes used by infected cells to elongate telomeres (discussed below). Recent work by Tan et al. revealed that activation of DNA damage response is required to trigger a robust type I interferons response (IFNs) following mitochondrial DNA damage(66), it is thus highly plausible that viruses evolved to interfere with the activation of the DDR in order to counteract the activation of an efficient antiviral response in infected cells. This type of IFNs activation is different from the nuclear factor $\kappa\beta$ (NF- $\kappa\beta$)-dependent IFNs production that rely only on MRE11 and RAD50(18). Further studies will be required to investigate this possibility as well as the requirement of NSB1 for this process.

In germline, hematopoietic, stem and rapidly renewing cells, telomere elongation relies on the hTERT, a polymerase that catalyzes the extension of telomeric DNA repeats using RNA as template(67). While hTERT is negatively regulated in somatic cells, senescence is overcome in cancer cells either through the re-activation of the hTERT enzyme or by an alternative homology-directed mechanism called ALT(68). The HHV-6B genome contains conserved telomeric sequences that are required for viral integration(14). In this study, we show that HHV-6B integration is independent of NBS1 in ALT- cells while it is dependent on NBS1 in ALT+ cells. These findings are consistent with previous report showing that the telomerase complex is required for optimal HHV-6B integration(34) as well as with the role of NBS1 in ALT (69, 70). While PML is not required for the interaction of IE1 with NBS1 and the ability of IE1 to inhibit the phosphorylation of H2AX (this study), NBS1 is required for the assembly of functional ALT-associated PML bodies(71). These concomitant roles are in line with the absence of phenotype associated with NBS1-depletion in integration assay performed on PML^{-/-} ALT+ U2OS. Intriguingly, we previously report that PML KO also reduces integration in the ALT- HeLa cells, reinforcing the

hypothesis that PML plays ALT-independent role in this process(33). Further studies will be required to elucidate this function.

In line with previous findings showing that HHV-6B integration is not altered upon inhibition of RAD51(72, 73), we found that IE1 inhibits homology-driven repair processes, and that integration is independent of NBS1 in ALT- cell lines. Together, these observations argue against models where integration mechanisms rely on RAD51-dependent BIR or SSA(74). However, it is important to note that all homology-directed reporter assays used in this study rely on extensive DNA end resection following the induction of breakage by the nucleases I-SceI or Cas9, a process that is dependent on NBS1(75). Thereby, integration models where SSA or RAD51-independent BIR trigger integration following extensive accumulation of single-strand DNA generated at stalled replication fork are still plausible. One attractive model is that the integration of HHV-6B occurs during mitotic DNA synthesis (MiDAS), a RAD52-dependent BIR mechanism that is initiated upon replication fork stall that remain unresolved at the start of mitosis, a problem often observed at DNA locus that are hard to replicate such as telomeres(9, 23, 76). Such mechanism is NBS1- and RAD51-independent and is mediated by RAD52, POLD3 as well as the structure-specific nuclease MUS81-EME1. Alternatively, upon entry into a cell and before the viral genome circularizes (and before IE1 is expressed), the viral DNA can be perceived as broken DNA. Under such circumstances, the MRN complex would be recruited to the ends of the viral genome and initiate 3'→ 5' resections. The ssDNA ends of eroded telomere (no longer efficiently protected by the shelterin complex) could anneal to the near terminal telomeric sequence at the right end of the genome in a process analogous to an ALT mechanism described in yeast (reviewed in (23)). Once the entire viral genome is copied, the telomeric repeats at the left end of the genome would serve as template for telomerase or ALT mechanisms to regenerate a telomere of appropriate length(77).

In conclusion, we provide a detailed characterization of the HHV-6B IE1 protein as an efficient inhibitor of DSB-signaling through the recruitment of NBS1. As such, IE1 contributes to the favorable establishment of a productive infection. Despite being a relatively abundant protein expressed very early upon entry, the functions of IE1 remain poorly defined. IE1 shares very little sequence homology with proteins from other herpesviruses (except HHV-6A and HHV-7) meaning that deductions based on primary sequence analysis are very limited. Our work adds to the growing knowledge surrounding HHV-6B integration processes and the potential importance of the IE1 protein during the infectious process.

Material and Methods

Plasmids and virus

pcDNA4/TO/myc-His-HHV-6B IE1 was previously described (27). The PiggyBac transposon-based (PB)-TetO and the PB-CA-rtTA-IRES-NEO plasmids were generated as previously described(78). PB-TetO-HHV-6B IE1, mCherry-LacR and GFP expression vectors were generated using Invitrogen™ Gateway™ recombination cloning (Invitrogen) and the following destination vectors: pDEST-PB-TetO (78), pDEST-mCherry-LacR (79) or pDEST-FRT-TO-GFP (80). HHV-6B IE1 was PCR amplified from pcDNA4/TO-HHV-6B IE1. HHV-6B IE1 fragments (aa1-1078, aa1-809, aa1-540, aa541-809, and aa541-1078) and NBS1 fragments (aa1-754, aa1-733, aa1-638, aa1-327, aa109-754, aa201-754, and aa328-754) were PCR amplified from pcDNA4/TO-HHV-

6B IE1 and pLXIN2-NBS1, a kind gift from Cary A. Moody (University of North Carolina at Chapel Hill, North-Carolina) (81). HHV-6B strain Z29 (82) was produced by our laboratory, as previously described(83). A list of the plasmids that were used in this study is provided in Table S3.

RNA interference

SMARTPool siRNA targeting RAD51 and a non-targeting single siRNA duplex sequences were purchased from Dharmacon. Single siRNA duplexes targeting RAD52 was a kind gift from Jean-Yves Masson (Université Laval, Québec, Canada). siRNAs were transfected in a forward transfection mode 24 hours prior to cell processing using RNAimax (Invitrogen) according to the manufacturer's protocol. Plasmids carrying a NBS1 short hairpin RNA (shNBS1) (TRCN0000010393, Open Biosystems) or a control shRNA (shCTRL) (Mission® TRC2 pLKO.5-puro non-mammalian shRNA control plasmid DNA, Sigma #SHC202) in the pLKO background backbone were a kind gift from Cary A. Moody (University of North Carolina at Chapel Hill, North-Carolina) (81). Lentiviruses were produced as previously described (81). Briefly, plasmids expressing shRNAs with vesicular stomatitis virus G (pMD2.g) and lentiviral packaging (pPAX) plasmids were co-transfected into HEK-293T cells using polyethyleneimine (PEI). 48-72 hours post-transfection, supernatants containing lentivirus were harvested and U2OS, MOLT-3, HeLa, and GM847 were transduced in the presence of 8 µg/ml hexadimethrine bromide (Polybrene) (Sigma). Knockdown of RAD51, RAD52 and NBS1 were confirmed for each experiment by Western blotting or qPCR analyses.

Cell Culture and transfections

Cell lines were maintained at 37°C and 5% CO₂. All culture media were supplemented with 10% fetal bovine serum (FBS). MOLT-3 (American Type Culture Collection, ATCC) were cultured in Roswell Park Memorial Institute (RPMI-1640; Corning Cellgro), 8.85 mM HEPES and 5 µg/ml plasmocin (Invivogen). GM847 and HeLa cell lines were obtained from ATCC and cultured in Dulbecco's Modified Eagle's Medium (DMEM; Corning Cellgro), NEM (Corning Cellgro), 8.85 mM HEPES, and 5 µg/ml plasmocin (Invivogen). U2OS (U2OS, obtained from ATCC), U2OS PML^{-/-} (37), U2OS 2-6-5 (Gift from Roger Greenberg, University of Pennsylvania, Philadelphia)(39), U2OS DR-GFP, NHEJ-GFP (EJ7), and SA-GFP (Gift of Jeremy Stark (City of Hope National Medical Center, California)(84, 85), and U2OS NHEJ-pc222 (Gift from Jacques Côté (Université Laval, Québec)(86) cell lines were cultured in McCoy's medium (Life Technologies). Doxycycline-inducible U2OS HHV-6B IE1 clones 10 and 102 (C10 and C102) were established by co-transfecting PB-TetO-HHV-6B IE1, pCMV-hypB_{ase} and PB-CA-rtTA-IRES-NEO plasmids, at a DNA ratio of 1:1:1 in the U2OS SA-GFP cell line using Lipofectamine 2000 (Invitrogen) according to the manufacturer's protocol. Clones were selected using 40 mg/mL of G418 and isolated using a limit dilution approach. U2OS BIR were established with a GFP-based reporter plasmid (pBIR-GFP) containing already characterized I-Sce1 reporter cassette to monitor BIR(87). Plasmid transfection was carried out using Lipofectamine 2000 (Invitrogen) according to the manufacturer's protocol. Clones were selected using 2 µg/ml puromycin and isolated using a limit dilution approach. Experiments were performed with a stable reporter clone which produce between 1.5% and 3% of GFP-positive cells after DSBs induction. Unless indicated otherwise, expression of IE1 was induced by adding 1 µg/ml doxycycline for 48h. HeLa DR-GFP (Gift from Roger Greenberg, University of Pennsylvania, Philadelphia) and HEK293T cell lines were cultured in Dulbecco's modified Eagle's medium (DMEM) (Life Technologies). All cell

lines were validated using short tandem repeat (STR) markers and tested negative for mycoplasma contamination.

Chemicals and sources of DNA damage

Doxycycline (Dox, 1 µg/ml, D3447, Sigma) was used to induce the production of HHV-6B IE1 in stable U2OS cell lines C10 and C102 for 48h. In the FOK1 system, DSBs were created at the LacO array by promoting the nuclear localization (4-Hydroxytamoxifen (4-OHT, 100 nM, #3412, Tocris)) and stabilization (Shield-1 ligand, 0.5 µM, CIP-S1-0001, CheminPharma)) of mCherry-LacR-FOK1 nuclease fused to a destabilization domain (DD) and to a modified estradiol receptor (ER) (ER-mCherry-LacR-FOKI-DD) for six hours prior to immunofluorescence sample preparation. DNA damage were induced by exposing cells to ionizing irradiation (IR). U2OS were exposed to 1 Gy with a CellRad (Precision X-Ray Inc.) and MOLT-3 to 4 Gy with the Gammacell® 40 Exactor (Best Theratronics Ltd.).

Viral infection and integration assays

For immunofluorescence assays on viral infection, 1 x 10⁶ MOLT-3 cells were pelleted, infected or not (Mock) at a multiplicity of infection (MOI) of 1 with HHV-6B (strain Z29) and resuspended at final concentration of 1 x 10⁷ cells/ml with fresh RPMI in a 1.5 ml tube for 5 hours at 37°C, 5%. The MOLT-3/HHV-6B prep was mixed every 30 minutes by flickering the tube. Cells were then washed three times in phosphate-buffered saline (PBS) and resuspended in 1 ml of fresh RPMI in a 12-well plate. Twenty-four hours post-infection, cells were processed for immunofluorescence. For viral replication, 1.5 x 10⁶ of MOLT-3 cells were pelleted into a 1.5 ml tube and infected or not (Mock) at a MOI of 1 with HHV-6B for 5 hours as described above. After 3 washes with PBS cells were resuspended in 3 ml of fresh RPMI, in a 6-well plate. At the indicated time point, 0.5 x 10⁶ cells were harvested and processed for DNA extraction using QIAamp DNA Blood Mini Kit as described by the manufacturer (Qiagen Inc.) and analysed by qPCR. Integration assays were performed as described previously(88). Briefly, 1 x 10⁴ cells/well (U2OS shCTRL, U2OS shNBS1, HeLa shCTRL, HeLa shNBS1, GM847 shCTRL, GM847 shNBS1), cells were infected at MOI of 1 with HHV-6B in a 24-wells plate for 24 hours and washed with PBS 1X 3 times. Cells were then seeded in 6-well plates and passaged for 4 weeks prior to DNA extraction with the QIAamp DNA Blood Mini Kit as described by the manufacturer (Qiagen Inc.) and analyzed by ddPCR.

Quantitative PCR (qPCR) and droplet digital PCR (ddPCR) analyses

qPCR was performed as previously described (83). DNA was analyzed using primers and probes against *U67-68* (HHV-6B) and *RPP30* (reference gene). Data were normalized against the corresponding genome copies of the cellular *RPP30* gene. ddPCR was used to quantify integration frequency as previously described (88). Briefly, the HHV-6B chromosomal integration frequencies were estimated assuming a single integrated HHV-6/cell and calculated with the following formula: (number of HHV-6 copies)/(number of RPP30 copies/2 copies per cell) × 100, as previously described. This assay was previously extensively validated and provide comparable data to single cell cloning and quantification.

RNA extraction and RT-qPCR

Total RNAs were extracted with the RNeasy mini kit following manufacturer's instructions (QIAGEN) and quantified by nanoDrop. 250-500 ng of total RNA was reverse transcribed with the High-Capacity cDNA reverse transcription kit (Invitrogen) according to the manufacturer's instructions. Contaminant genomic DNA was removed by DNaseI (ThermoFisher) incubation prior to the reverse transcription (RT) reaction and confirmed by GAPDH RT-PCR performed on DNaseI treated reactions. qPCR was performed using the LightCycler 480 apparatus (Roche) with the LightCycler 480 SYBR Green 1 qPCR master mix (Roche) using the following program: 40 cycles of 94 °C denaturation for 15 sec, 56 °C annealing for 5 sec and 72 °C elongation for 15 sec. 5% of the RT-PCR reaction was used as template. Standard curve was performed with serial dilution using the U2OS cDNA as template. Relative expression of each gene was determined using the standard curve and normalized to the relative expression of the GAPDH. The primers are listed in Table S1.

Immunofluorescence microscopy

One hour post-irradiation, MOLT-3 cells were pelleted, washed 3 times in PBS and 1×10^4 cells were added to each well of a microscope slide with 10 reaction wells (MP Biomedicals™ Multitest Slides, Fisher Scientific # ICN6041805). Once dried, cells were fixed for 20 minutes at room temperature with 2% paraformaldehyde (PFA), hydrated for 5 minutes with PSB and processed for immunofluorescence. U2OS and U2OS 2-6-5 cells were grown in 24-well plates on glass coverslips and fixed 24 hours later with either 2% (wt/vol) PFA in PBS for 20 minutes at room temperature or with 100% MeOH for 20 minutes at -20°C. When indicated, cells were treated with the indicated amount of Gy, 15 min or 1 hour prior to fixation. For immunostaining with anti-NBS1 and anti-MRE11 antibodies, nuclear extraction with ice-cold NuEx buffer (20 mM HEPES pH 7.4, 20 mM NaCl, 5 mM MgCl₂, 0.5% NP-40, protease inhibitors (Complete, EDTA-free protease inhibitor, Sigma), and 1 mM DTT) for 20 min on ice for prior to fixation. U2OS PFA-fixed cells were further permeabilized with 0.3% (vol/vol) Triton X-100 for 20 minutes at room temperature. MOLT-3 and U2OS fixed cells were incubated with blocking buffer (2% BSA in PBS or 0.1% BSA, 3% goat serum, 0.1% Triton, 1 mM EDTA pH 8.0 in PBS) for 30 minutes at room temperature and then incubated with primary antibodies (Table S2) diluted in blocking buffer for 2 hours at room temperature, followed by washes in PBS. Next, cells were incubated for 1 hour at room temperature with secondary antibodies diluted in blocking buffer and counterstained with 4',6-diamidino-2-phenylindole (DAPI, 0.4 µg/mL) in PBS. Cells were washed with PBS 1X and the coverslips were mounted onto glass slides with Prolong Diamond mounting agent (Invitrogen). To visualize micronuclei in MOLT-3 infected or control cells, cells were collected processed with an hypotonic solution (75 mM KCl) for 20 minutes at 37°C. Cell were then fixed with fresh 3:1 methanol:acetic acid for 5 minutes and washed three time with 3:1 methanol:acetic acid solution. Washed pellets were resuspended in 500µl of 3:1 methanol:acetic acid, dropped on a microscope slide and air dried prior to DNA counterstaining with DAPI. Images were either taken using a Zeiss LSM700 (and LSM900 recently acquired) laser-scanning microscope equipped with a 63x oil lens or a Wave FX-Borealis - Leica DMI 6000B microscope with the camera Image EM (Hamamatsu, 512x512 pixels) and Orca-R2 (Hamamatsu, 1344x1024 pixels) with a 40x (Quorum Technologies). Images were analyzed and quantified using ImageJ software [National Institutes of Health (NIH)]. In micrographs, dashed lines indicated nucleus outlines when DAPI is not shown. Unless stated otherwise, insets represent 10 X magnifications of the indicated fields.

***in situ* hybridization (FISH)**

Fixed cells were processed as described for immunofluorescence staining and then fixed for 2 minutes at room temperature with 1% PFA/PBS. Coverslips were washed twice with PBS for 5 minutes and dehydrated for 5 minutes in successive ethanol baths (70%, 95%, 100%). Once dried, coverslips were placed upside down on a drop of hybridizing solution (70% formamide; 0.5% blocking reagent; 10 mM Tris-HCl pH 7.2; 1/1000 Cy5-TelC PNA probe (F1003, PNABio, Newbury Park, CA, USA)). Sample were denatured for 10 minutes at 80°C on a heated block. Coverslips were incubated over night at 4°C and kept in the dark. After hybridization, coverslips were washed two times for 15 minutes in washing solution (70% formamide; 10 mM Tris-HCl pH 7.2) and then washed 3 times for 5 minutes with PBS. Sampled were air dried, counterstained with DAPI, washed with PBS and coverslips were mounted onto glass slides with Prolong Gold mounting agent (Invitrogen).

Metaphase spread analysis

U2OS SA-GFP HHV-6B IE1 cells were arrested in mitosis using 1 μ M nocodazole for 3 hours at 37°C and 5% CO₂. Cells were then resuspended and incubated in pre-warmed hypotonic solution (KCl 75 mM, 15% SVF) at 37°C for 15 minutes to induce swelling and fixed in (75% ethanol, 25% acetic acid) overnight at 4°C. Droplet of cells were spread onto glass slides pre-cooled at -20°C and dried overnight in the dark at room temperature. Slide were then mounted with Vectashield Antifade Mounting Medium containing DAPI (VECTH20002, MJS BioLynx Inc.). Images were taken using a Zeiss LSM700 laser-scanning microscope equipped with a 40x water lens. Quantification was done on 3 biological replicates and 10 spreads were quantified per experiments.

Reporter-based DNA repair assays

For DR-, NHEJ-, SA-, and BIR-GFP reporter assays in which HHV-6B IE1 was transiently transfected, U2OS or HeLa cells carrying the respective GFP expression cassette were plated at 125 000 cells/well in a 6 well plates. Twenty-four hours later, cells were co-transfected with the indicated combination of plasmids: pcDNA4/TO-HHV-6B IE1 along with I-SceI plasmid (pCBASceI, Addgene #26477). The pcDNA4/TO/myc-His vector was used as negative control for conditions without I-SceI or IE1. A plasmid expressing iRFP was also transfected to correct for transfection efficiency in each assay. For the NHEJ-GFP (EJ7) assay, cells were co-transfected with Cas9 and sgRNA-expressing vectors p330X-sgRNA7a and p330X-sgRNA7b expressing plasmids instead of I-SceI(85). After 48 hours (or 72 hours for NHEJ-GFP (EJ7)), cells were trypsinized, harvested, washed and re-suspended in PBS. The percentage of GFP-positive in iRFP-positive cells was determined by flow cytometry using an Accuri C6 (BD Biosciences). The data were analyzed using the FlowJo software (Flow Jo LLC). When indicated, cells were transfected with siRNA 24 hours prior to transfection with I-Sce1 expression plasmid. For SA-GFP and the CRISPR-LMNA HDR assay that were done in U2OS SA-GFP HHV-6B IE1, cells were seeded at 10 000 cells per well in 24-well plates and induced with 1 μ g/mL of doxycycline 24 hours post transfection with either I-Sce1 for SA-GFP assay or plasmid expressing Cas9 and LMNA sgRNA (pX330-LMNAsgRNA1) and CR2.1-mRuby-2-LMNA-Donor for CRISPR-LMNA HDR assay(43). At 48 hours post-transfection, cells were harvested and GFP-positive cells quantified by flow cytometry. mRuby-positive cells were analyzed by microscopy using a TIRF Ti-LAPP microscope (Nikon).

Statistical analysis

Quantification was done on 3 biological replicates. Unless stated otherwise, one-way ANOVA with Dunnett's multiple comparisons test were realized to assess statistical significance.

Acknowledgement

We are grateful to Matthew D. Weitzman, Alexandre Orthwein, Cary A. Moody and members of the A.F.-T. and L.F. laboratory for critical reading of the manuscript; and to Daniel Durocher, Jean-Yves Masson, Graham Dellaire, Roger Greenberg, and Jeremy Stark for essential reagents such as plasmids, purified proteins, and cell lines. E.B. and V.C. received a postdoctoral and a doctoral fellowship from the Fonds de Recherche du Québec - Santé (FRQS), respectively. V.T. received a master fellowship from the Fonds de recherche Nature et technologies (FQRNT). This work was supported by three Canadian Institutes of Health Research Grants (PDT_152948 to A.F.-T., and MOP_123214 and PJT_156118 to L.F.). A.F.-T. is a tier 2 Canada Research Chair in Molecular Virology and Genomic Instability and is supported by the Foundation J.-Louis Lévesque. We thank the Bioimaging platform of the Infectious Disease Research Centre, funded by an equipment and infrastructure grants from the Canadian Foundation Innovation (CFI).

Author Contributions: V.C., E.B., L.F. and A.F.-T. designed research; V.C., E.B., V.T., E.G.L., J.D. and A.G. performed research and analyzed data; V.C. and A.F.-T. wrote the original draft and E.B., and L.F. edited the manuscript.

Competing Interest Statement: The authors declare no conflict of interest.

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Figure Legends

Fig. 1. HHV-6B infection and IE1 expression leads to micronuclei formation. (A) Left panel: Representative images of micronuclei observed in HHV-6B infected MOLT-3 cells. Cells were infected and fixed 24 hours post-

infection. DNA was counterstained with DAPI. Micronuclei are indicated by white arrows (scale bar, 5 μ m). Quantification of micronuclei are presented on the right panel. Data are presented as the mean ($n = 2$, >100 micronuclei/condition). (B) Left panel: representative images of U2OS cell line and U2OS clones stably expressing doxycycline (Dox)-inducible HHV-6B IE1 protein (C10 and C102). Expression of IE1 was induced for 48 hours with 1 μ g/ml of Dox prior to fixation. Cells were then processed for IE1 immunofluorescence and counterstained with DAPI. Micronuclei are indicated by white arrows (scale bar, 5 μ m). Quantification of micronuclei are presented on the right panel. The parental cell line (Par.) was used as a negative control and data are presented as the mean $\pm$ SD ($n = 3$). (C) Schematic representation of the mechanisms leading to micronuclei formation. Events leading to the formation of micronuclei induced by DNA double-strand breaks (DSBs) (i) and lagging chromosome (ii) or by anaphase bridges (ABs) (iii) are represented. (D-E) Quantification of micronuclei containing centromere (D) and telomere (E). Cells were treated as described in B and either processed for centromere immunofluorescence or by FISH for the detection of telomeres. Data are presented as mean $\pm$ SD ($n = 3$) (D) and as the mean ($n = 2$, >100 micronuclei/condition) (E). (F) Representative image of a metaphase from IE1 expressing cells. Cells were exposed to 1 μ g/ml of dox for 48 hours, metaphase spread were prepared, fixed and processed for DNA counterstaining. (G) Quantification of chromosomal aberrations per metaphase. Data were analyzed with an unpaired t-test and are presented as mean $\pm$ SD ($n = 31$). ** $p < 0.01$, *** $p < 0.001$, **** $p < 0.0001$.

Fig. 2. Phosphorylation of H2AX (γ -H2AX) is inhibited in HHV-6B infected and IE1 expressing cells. (A) Representative images of γ -H2AX in irradiated U2OS parental (Par.) and IE1-expressing cells. Cells were treated as described in Fig. 1B and irradiated with 1 Gy. One hour post-irradiation, cells were fixed and processed for IE1 and γ -H2AX immunofluorescence (scale bar, 5 μ m). (B) Quantification of cells with more than 10 γ -H2AX foci in irradiated U2OS Par. and IE1-expressing cells. Data are presented as the mean $\pm$ SD ($n = 3$). (C) Quantification of cells with more than 10 γ -H2AX foci in U2OS PML^{+/+} and ^{-/-} irradiated cells (1 Gy) that transiently express untagged IE1. An empty vector (EV) was used as negative control. Data are presented as the mean $\pm$ SD ($n = 3$). (D) Representative images of γ -H2AX in HHV-6B infected MOLT-3 cell lines. Cells were irradiated with 4 Gy. One hour post-irradiation, cells were fixed and processed for IE1 and γ -H2AX immunofluorescence (scale bar, 5 μ m). Mock-infected cells were used as a negative control. (E) Quantification of cells with more than 10 γ -H2AX foci in irradiated MOLT-3 infected cells. Data are presented as the mean $\pm$ SD ($n = 3$) and statistical significance was assessed using unpaired t-test. **** $p < 0.0001$.

Fig. 3. HHV-6B IE1 interacts with NBS1 and prevents its recruitment to DSBs. (A, C) Representative images of the colocalization between NBS1 (A) and MRE11 (C) with IE1. IE1-expressing cells were treated as described in Fig. 1B, fixed and processed for IE1, NBS1, or MRE11 immunofluorescence as indicated. As a positive control, irradiated U2OS cells (+IR) were fixed 15 minutes post-irradiation (1 Gy) and processed for γ -H2AX, NBS1, or MRE11 immunofluorescence as indicated (scale bar, 5 μ m). The parental cell line (Par.) was used as a negative control. (B, D) Quantification of γ -H2AX or IE1 foci that colocalized with NBS1 (B) and MRE11 (D) are presented as percentage of foci per cells that colocalized with the indicated protein. Data are presented as the mean $\pm$ SD of three independent experiments. (E) U2OS 2-6-5 cells transfected with plasmids expressing

the indicated mCherry-LacR fusion protein or induced for the expression of ER-mCherry-LacR-FokI-DD were fixed and processed for NBS1 immunofluorescence (scale bars, 5 μ m). The mCherry-LacR backbone was used as a negative control (--). (F) Quantification of the mCherry-LacR foci colocalizing with NBS1 (E), γ -H2AX (SI Appendix, Fig. S3F), and MRE11 (SI Appendix, Fig. S3G). Data are presented as the mean $\pm$ SD ($n = 3$). **** $p < 0.0001$.

Fig. 4. Bipartite binding and inhibition of NBS1 by HHV-6B IE1. (A) Schematic representation of HHV-6B IE1 protein and the fragments of the protein used in this study. NID, NBS1-interacting domain; NBS1i, NBS1 inhibitory domain, STAT2-BD: STAT2 binding-domain (aa 270-540). S432: CDK2 phosphorylation site. (B) U2OS 2-6-5 cells transfected with the plasmids expressing the indicated mCherry-LacR fusion protein or induced for the expression of ER-mCherry-LacR-FokI-DD were fixed and processed for NBS1 immunofluorescence (SI Appendix, Fig. S4A-B). The mCherry-LacR backbone was used as a negative control (--). (C) Representative images of the inhibition of γ -H2AX by IE1. Cells were transiently transfected with the indicated mCherry-LacR fusion protein and irradiated 24 hours later. One hour post-irradiation (1 Gy), cells were fixed and processed for γ -H2AX immunofluorescence. The mCherry-LacR backbone was used as a negative control (--). (D) Quantification of cells with more than 10 γ -H2AX foci. UT, untreated. Data for (B) and (D) are presented as the mean $\pm$ SD ($n = 3$). (E) U2OS 2-6-5 cells were treated as described in (B), processed for IE1 and γ -H2AX immunofluorescence (SI Appendix, Fig. S4E-F) and quantified as indicated. (F) Schematic representation of NBS1 protein and the fragments of the protein used in this study. FHA, ForkHead-Associated domain; BRCT, BRCA1 C-Terminal domain; MRE11-BM, MRE11-binding motif; ATM-BM, ATM-binding motif; IDD, Intrinsically Disorder Domain. (F-G) U2OS 2-6-5 cells transfected with the plasmids expressing the indicated mCherry-LacR and were fixed and processed. Data for (E) and (G) are presented as the median $\pm$ SD ($n = 3$) (E). ** $p < 0.01$, **** $p < 0.0001$.

Fig. 5. HHV-6B IE1 inhibits HDR-mediated repair. DNA repair reporter assays for (A) homologous recombination (DR-GFP), (B) Single-strand annealing (SA-GFP), (C) Break-induced replication (BIR-GFP) and (D) Non-homologous end-joining (NHEJ-GFP (EJ7)). For each condition, a schematic representation of the assay is presented in the top panel and quantification of the GFP+ cells analyzed by flow cytometry is presented in the bottom panel. GFP-positive cells are normalized over GFP-positive cells quantified in the positive control (I-SceI+, set to 1.0) in each replicate. Data are represented as the mean $\pm$ SD ($n = 3$). **** $p < 0.0001$.

Fig. 6. Depletion of NBS1 impairs viral integration in cells maintaining telomere by homology-directed repair. (A) Schematic representation of HHV-6B infection in permissive and semi-permissive cells. In semi-permissive cells for HHV-6B, where replication is inefficient, and the viral genome integrates at telomeres. (B) MOLT-3 cells depleted or not for NBS1 were infected at a MOI of 1 with HHV-6B and harvested at the indicated time points. Following cell lysis, DNA was extracted and the number of copies of HHV-6B were determined by qPCR using primers for U67-68 gene for HHV- 6B and RPP30 as a cellular reference gene. Data presented are the mean of three independent experiment and presented as mean $\pm$ SD ($n = 3$).

Supplementary Figure Legends

Fig. S1. (A-B) Whole cell extracts (WCE) from infected MOLT-3 (A) and U2OS cells treated or not with 1 μ g/ml of Dox (B) were analyzed by immunoblotting with an antibody against IE1. β -tubulin was used as loading control. NI: non-infected, Par.: parental cell line. (C, F, G) Representative images of the U2OS (Par.) and U2OS IE1 stable cell lines with or without Dox induction (as indicated) for Figure 1B, D and E. Cells were treated as described in Fig. 1B and either processed for IE1 (C, G) or centromeres (CREST) immunofluorescence (F) or by FISH for the detection of telomeres (G) (scale bar, 5 μ m). (D-E) U2OS (Par.) and U2OS IE1 stable cell lines with Dox induction were treated as described in Fig. 1B and processed for Lamin B fluorescence. Quantification of micronuclei with Lamin B signal is presented in (D) and representative images in (E). In (D), data are represented as mean $\pm$ SD ($n = 3$) ** $p \leq 0.01$. (H) Quantification of micronuclei colocalizing with IE1 foci. IE1-expressing cells were treated as described in Fig. 1C, fixed and processed for IE1 immunofluorescence. Data are presented as the mean $\pm$ SD ($n = 2$, >100 nuclei/condition).

Fig. S2. (A) Representative images of untreated U2OS PML^{+/+} and ^{-/-} cells fixed and processed for PML immunofluorescence. (B) Western blot analysis of U2OS transfected with untagged IE1 or an empty vector (EV) plasmids WCE. β -tubulin was used as loading control. (C) Representative images of U2OS PML^{+/+} and ^{-/-} cells transiently transfected with a plasmid expressing untagged IE1 or an empty vector (EV) as negative control. Cells were irradiated with 1 Gy, fixed 15 minutes post-irradiation and processed for IE1 and γ -H2AX immunofluorescence (scale bar, 5 μ m).

Fig. S3. (A) WCE from U2OS cells (Par.) and IE1-expressing U2OS stable cell lines treated or not with 1 μ g/ml of Dox were analyzed by immunoblotting with antibodies against RAD50, NBS1 and MRE11. GAPDH was used as loading control. (B-C) Representative immunofluorescence of IE1-expressing U2OS stable cell lines induced with Dox and irradiated with 1 Gy. Cells were fixed 1 hour post-irradiation and processed as described in Fig. 3A and C, respectively. (D) Quantification of IE1 foci that colocalize MRE11 in stable U2OS control cells (shCTRL) or depleted for NBS1 (shNBS1). Cells were transiently transfected with untagged IE1 and treated as described in Fig. 3B. Percentage of IE1 foci per cells that colocalize with NBS1 are presented as the mean $\pm$ SD ($n = 2$, at least 40 nuclei/condition). (E) Representative immunofluorescence of the data presented in Fig. Supp. 3D. (F) WCE from U2OS shCTRL and shNBS1 stable cell lines were analyzed by immunoblotting with antibodies against NBS1 and MRE11. GAPDH was used as loading control. (G-H) Representative immunofluorescence of the quantification presented in Fig. 3F. (I) Quantification of IE1 foci that colocalize with NBS1 in U2OS PML^{+/+} and ^{-/-} cells transiently expressing untagged IE1. Cells were treated as described in Fig. 3A. Percentage of IE1 foci per cells that colocalize with NBS1 are presented as the mean $\pm$ SD ($n = 3$). (J) Representative immunofluorescence of the quantification presented in SI Appendix, Fig. S3I. Statistical significance in (D) was assessed by unpaired t-tests, **** $p < 0.0001$.

Fig. S4. (A, D) WCE from U2OS cells that transiently express the indicated mCherry-LacR fusion protein were analyzed by immunoblotting with antibodies against mCherry. GAPDH was used as loading control. The mCherry-LacR backbone was used as a negative control (--) in (A). (B-C) Representative immunofluorescence

images used for the quantification presented in Fig. 4B (B), Fig. 4D (C) (scale bar, 5 μ m). (E) Representative immunofluorescence images used for the quantification presented in Fig. 4E and Fig. Supp 4F (scale bar, 5 μ m). (F) Quantification of mCherry-LacR foci dans colocalize with IE1. Data are presented as the median $\pm$ SD ($n = 3$) **** $p < 0.0001$. (G) Representative immunofluorescence images used for the quantification presented in Fig. 4G and Fig. Supp 4H (scale bar, 5 μ m)

Fig. S5 (A) CRISPR-LMNA HDR assay was analyzed in Dox-inducible IE1 U2OS SA-GFP stable cell lines. Cell lines were plated and induced for IE1 expression for 24 hours prior to transfection with plasmids encoding Cas9, LMNA sgRNA, and mRuby2-LMNA donor. Percentage of mRuby-positive cells were analyzed by flow cytometry 48 h post-transfection and normalized over percentage of mRuby-positive U2OS SA cells (Par.) in each replicate. (B) DNA repair assay for homologous recombination in HeLa cells were performed as described for U2OS cells in Fig. 5A. (C) Single-strand annealing (SA) assay was analyzed in Dox-inducible IE1 U2OS SA-GFP stable cell lines. Cell lines were plated and induced for IE1 expression for 24 hours prior to transfection with plasmids encoding I-SceI endonuclease. Percentage of GFP-positive cells were analyzed by flow cytometry 48 h post-transfection and normalized over percentage of GFP-positive U2OS SA cells (Par.) in each replicate. Data are represented as the mean $\pm$ SD ($n = 2$). (D) Validation of the BIR repair assay using siRNA against RAD51 and RAD52. A non-targeting siRNA (NT) was also used as control. (E-F) RT-qPCR was performed on U2OS BIR cells using gene-specific primers for RAD51(E) and RAD52(F). Expression of each transcript has been normalized against GADPH. (G) DNA repair assay for non-homologous end-joining (NHEJ-pc222) in U2OS cells were performed as described in Fig. 5D. Unless stated otherwise, data are represented as the mean $\pm$ SD (at least $n = 3$). In (A), statistical significance was assessed by unpaired t-tests, ** $p < 0.01$, *** $p < 0.001$, **** $p < 0.0001$.

Fig. S6. (A-D) WCE from MOLT-3 (A), HeLa (B), U2OS (C), and GM847 (D) cell lines expressing a shRNA against NBS1 (shNBS1) or control (shCTRL) were analyzed by immunoblotting with an antibody against NBS1. β -Tubulin was used as loading control.

Fig. 1

Fig. 2

FIGURE 3

Fig. 5

Fig. 6

A

HHV-6B infection

Permissive cell

Lytic state
Viral replication

Semi-permissive cell

Latent state
Viral integration

Viral
episome

Human telomere
DNA

Linear viral DNA

B