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The Epstein-Barr virus (EBV) protein EB2 is an mRNA export factor essential for virus production

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1. ABSTRACT

The EBV early protein EB2 (aka Mta, SM and BMLF1) shares properties with mRNA export factors. It shuttles between the cytoplasm and the nucleus, and interacts with RNA both *in vitro* and *in vivo* but with no apparent sequence specificity. EB2 induces the cytoplasmic accumulation of mRNAs generated from intronless and intron-containing genes, likely through interactions with cellular export factors of the TAP/p15 pathway. Using a cell line carrying a viral genome with the EB2 gene deleted, it has been shown that EB2 is essential for the production of infectious virions by facilitating the nuclear export of a subset of early and late viral mRNAs, a function regulated by CK2 phosphorylation of EB2. There are docking sites for both CK2 subunits and for the heterotetrameric enzyme in the EB2 N- and C-terminal domains. Accordingly, EB2 and CK2 co-purify as a complex in which CK2 phosphorylates EB2. CK2 phosphorylation of EB2 at one of the Ser-55, Ser-56 and Ser-57 is critical for its mRNA export function and as a consequence, for infectious virus production.

2. INTRODUCTION

The Epstein-Barr virus infects more than 95% of the human adult population. The primary infection is generally silent, but can be the direct cause of a benign disease called infectious mononucleosis (1). After the primary infection, EBV persists lifelong in a latent state, in a sub-population of resting memory B cells (2). The persistence is considered a risk factor for the emergence of EBV-associated malignancies Burkitt's Lymphoma (BL), Nasopharyngeal Carcinoma (NPC), gastric carcinomas, Hodgkin's disease, post-transplant lymphoproliferation disorders (PTLD), T/NK nasal lymphomas, AIDS-associated lymphomas and leiomyosarcomas (reviewed in (3, 4)). EBV has also recently been found in some breast carcinomas biopsies (5-7). Although the hallmark of latency is the absence of a complete viral productive cycle *in vitro*, EBV productively replicates *in vivo* in mucosal epithelia and infectious virions are found intermittently in the saliva of healthy carriers (8). This is likely due to the latent infection of blood-borne mononuclear cells that migrate to the oral epithelium where they differentiate into

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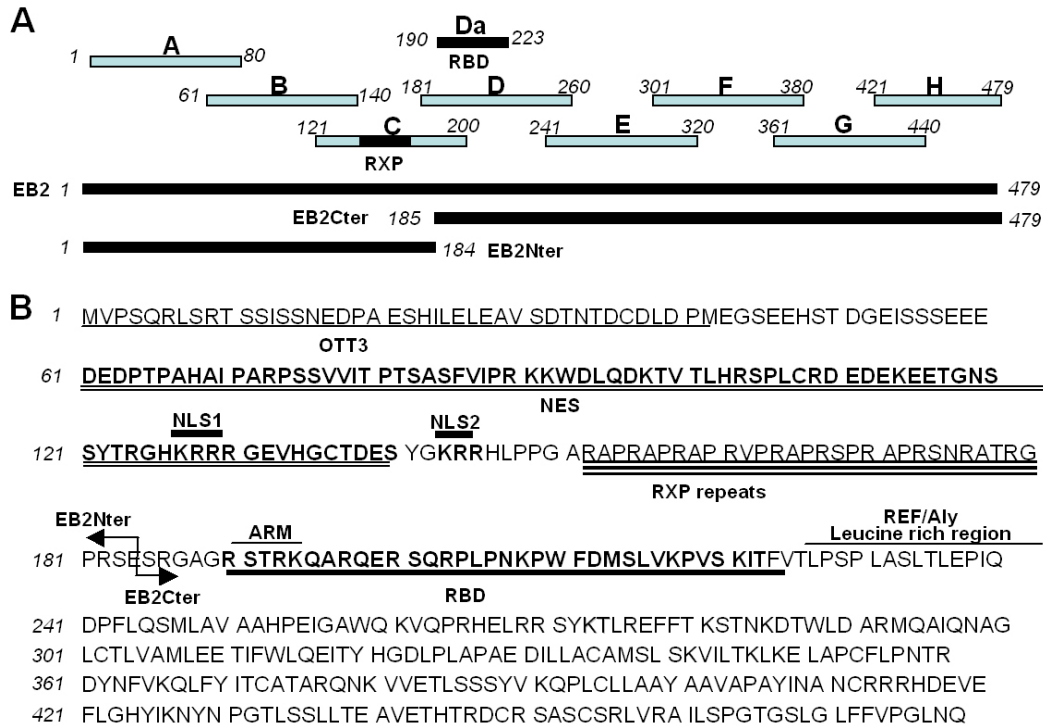


Figure 2. EB2 functional domains and interacting factors. A)- EB2 overlapping polypeptides that were fused to the C-terminus of GST to identify functional domains and interacting proteins. RXP: Arginine-X-Proline repeats; RBD: RNA Binding Domain. B)- Functional domains and interacting proteins are located on the EB2 amino acid sequence. The double arrow delineates the N-terminal fragment of EB2 (EBNter) and the C-terminal fragment of EB2 (EB2Cter). NES: Nuclear Export Signal; NLS: Nuclear Localization signal; ARM: Arginine-Rich Motif.

termination codon, at position 70455 (Figure 1). From this spliced mRNA, a 479 amino-acid polypeptide is expressed, generating the EB2 protein. The BMLF1 ORF putative protein product should not be expressed from the two small mRNAs unless there is an internal ribosomal entry site between the BSLF2 and the BMLF1 ORFs and this has not as yet been established. Nevertheless, in one report, co-expression of two proteins, one initiated at the BSLF2 AUG and the other at the BMLF1 AUG have been described in EBV-infected B cells (24). However, the protein initiated at the BSLF2 AUG was much more abundant than the protein initiated at the BMLF1 AUG. No other report has mentioned the expression of the shorter BMLF1 protein. The longer mRNAs, which also contain the BSLF1 ORF, have not been further characterized, and it is not known if EB2 is expressed from these mRNAs. The EB2 gene is an early gene and in the pM promoter a proximal AP-1 site, which is a TPA- and EB1-Responsive Element (ZRE) (27), and a distal R-Responsive Element (RRE) (28) have been characterized (Figure 1). Accordingly, the EB2 gene is transcribed under the control of both EB1/ZEBRA and R. The upstream pM1 promoter has not yet been characterized

4. EB2 SHARES PROPERTIES WITH mRNA EXPORT FACTORS

The EB2 protein was first described as a nuclear protein and a transactivator of gene expression as in

transiently transfected cells it activated expression of the chloramphenicol acetyl transferase (CAT) intronless gene placed under the control of many different promoters (19, 26, 29, 30). However, it was later shown that the effect of EB2 on gene expression was not promoter dependent but reporter gene dependent and was probably post-transcriptional (27, 31). It was finally demonstrated that EB2 has the properties of an RNA export factor and increases the cytoplasmic accumulation of certain intronless viral mRNAs in transient expression assays (32).

4.1. EB2 shuttles between the nucleus and the cytoplasm

Most RNA export factors shuttle between the nucleus and the cytoplasm and shuttling is essential for their mRNA export function. Shuttling is dependent on two short amino-acid sequences: a Nuclear Localization Signal (NLS) and a Nuclear Export Signal (NES). Transfer of RNA export factors between both compartments is mediated by soluble receptors that bind to the NLS and NES and interact with components of the nuclear pore complex (NPC) (reviewed in (33)). There are two NLSs in EB2, NLS1 and NLS2 (Figure 2B), which act independently and can be transferred to heterologous proteins (34). The cellular receptors that bind these NLS and facilitate the nuclear import of EB2 are not known as yet.

For the prototype RNA export factor Rev from HIV, the nuclear export signal (35) is leucine-rich and

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interacts with CRM1 in a complex including ranGTP, a GTPase in its GTP-bound state (36-39). CRM1 directs Rev to the nuclear pore complex for export (reviewed in (40)). A mutation in the leucine-rich NES of Rev impaired both Rev shuttling and Rev-dependent export of unspliced viral mRNAs (41). EB2 was at first also reported as being CRM1-dependent for its cytoplasmic translocation and to directly interact with CRM1 via a leucine-rich region (Figure 2B) (42). It was further shown that this leucine-rich region contains two putative NESs that when individually fused to a fluorescent nuclear reporter protein induced its cytoplasmic translocation in a CRM1-dependent manner (43). However, other studies using heterokaryon assays, reported that EB2 shuttling was CRM1-independent (44). Indeed, the antibiotic leptomycin B (LMB), a specific CRM1 inhibitor (45), impaired Rev- but not EB2-nucleocytoplasmic shuttling. Furthermore, replacement of the leucines by alanines in the EB2 leucine-rich region, a mutation that abolishes the activity of the CRM1-dependent leucine-rich NES (41), had no effect on the nucleocytoplasmic shuttling of EB2 (44). Deletion of the leucine-rich region also did not affect EB2's shuttling (34). The final demonstration that the leucine-rich region was not a NES came from microinjection into single nuclei of HeLa cell polykaryons, of the fusion proteins made between the Glutathione-S-Transferase (GST) and the EB2 overlapping peptides A to H (Figure 2A). In these assays, GST-B was transferred to all nuclei of the injected polykaryons, indicating that peptide B contains both an NLS and a NES (Figure 2B), while GST-D which contains the leucine-rich region, remained in the injected nuclei (46). The EB2 N-terminal NES, contained in the B peptide, was transferable and conferred CRM1 independent nucleocytoplasmic shuttling to beta-galactosidase. However, although EB2 shuttles between the cytoplasm and the nucleus, it localizes essentially to the nucleus, suggesting that the EB2 import rate is greater than the export rate when substrate mRNAs are limiting. This was also observed for ICP27, the HSV-1 homologue of EB2 in a study using *Xenopus laevis* oocytes (47). It is noteworthy that deletion of the B region in EB2 (targeted to the nucleus by a heterologous NLS in order to compensate for the concomitant deletion of one of the two EB2 NLS) did not impair nucleocytoplasmic shuttling, suggesting that there is a second NES in EB2 (our unpublished results). The second NES is evidently not the leucine-rich region since the C-terminal part of EB2 (Figure 2B, EB2Cter), targeted to the nucleus by a heterologous NLS, does not shuttle, even though it contains the leucine-rich region (34). On the contrary, the N-terminal part of EB2 (Figure 2B, EB2Nter) which contains the CRM1-independent NES does shuttle (our unpublished results). Thus, there are probably more than one NES in EB2, all localized in the N-terminal part of the protein. It is not yet known which cellular factor(s) binds to these N-terminal NESs of EB2 to facilitate its nuclear export.

4.2 EB2 interacts with RNA *in vitro* and *in vivo*

RNA export factors can bind RNA via a domain that directly recognizes either a specific RNA sequence (the case for some members of the family of splicing factors containing serine-arginine repeats (SR proteins)) (48) or a specific RNA structure (as in the case for TAP/NXF1 (49),

HIV Rev (50) and RBM15/OTT1 (51)). RNA export factors can also be recruited to RNA via interaction with components of complexes loaded onto RNA during splicing (52, 53), transcription elongation (54) or 5' capping (55).

By using the GST fusion peptides depicted in Figure 2A, in RNA mobility shift assays, it has been shown that EB2 contains an RNA-binding domain (RBD) located in peptide D (34). The EB2 RBD contains the motif TRKQAR, rich in basic residues (Figure 2B, ARM: Arginine Rich Motif), which is reminiscent of the arginine-rich motif TRRRER originally found in the bacteriophage lambda N protein (56) and then in many other RNA-binding proteins including HIV-1 Rev (TRQARR) and Tat (TRGKGR) (57, 58). The shorter RBD-containing peptide Da (Figure 2A and Figure 2B), fused to GST, bound *in vitro* with no apparent specificity to 180 and 72 nucleotide RNAs corresponding respectively to the second exon and the first intron of the human beta-globin gene (34). GST-Da bound several times to both RNA molecules, and the number of GST-Da:RNA complexes increased with increase in GST-Da concentration. Although the RNAs used in the *in vitro* binding assays were potentially highly structured as evaluated by MFOLD (59), GST-Da bound to both RNAs every 20 nt or so. Purified full length EB2 also bound *in vitro* several times to both RNAs (our unpublished data). It would thus appear that the RNA-binding domain of EB2 can directly interact with RNA *in vitro* and occupies sites covering the entire RNA molecule, suggesting that the EB2 RBD could unfold the RNA *in vitro*.

Another RNA-binding domain has been described for EB2 which contains several Arginine-X-Proline repeats (Figure 2B, RXP repeats), similar to that present in the HSV-1 US11 protein (60). The RXP repeats bound RNA *in vitro* when exposed at the C-terminus of the GST-RXP fusion protein (61), or after proteolytic cleavage of the GST-EB2 fusion protein and exposure at the C-terminus of the truncated protein (62). A shorter protein initiated at the BMLF1 AUG and fused at the C-terminus of GST also bound RNA *in vitro* but again the fusion protein was degraded and the RXP domain was probably exposed at the C-terminus of the GST-BMLF1 fusion protein (32). However, the RXP repeats did not bind RNA when embedded within the GST-C fusion protein and EB2Cter which lacks the RXP repeats (Figure 2A), bound RNA in a Northwestern assay (34). Furthermore, EB2 could be co-immunoprecipitated with target RNAs from transfected cells when the RXP repeats were deleted (62) but not when the RBD was deleted (34), strongly indicating that EB2 directly interacts with RNA in living cells via the RBD. Again no specific RNA sequence or structure was identified for directing EB2 to target RNAs *in vivo*.

4.3 EB2 interacts with cellular RNA export factors

The assembly of the export-competent mRNPs in mammalian cells, starts with capping at the 5'-end of the nascent RNA and subsequent binding of the cap-binding complex composed of CBP80 and CBP20. This step is followed by splicing-dependent deposition of the exon-

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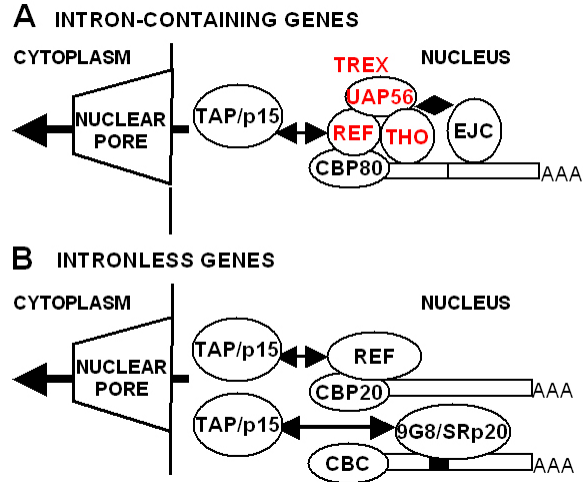


Figure 3. Models for cellular mRNA export. A)- Nuclear export of mRNAs that undergo splicing. 5' recruitment by CBP80 of REF/Aly bound to UAP56 in the TREX complex is splicing dependent, probably via interaction with the Exon Junction Complex (EJC), which is deposited on the RNA during splicing. TAP/p15 facilitates the translocation through the nuclear pore of the export-competent mRNP by contacting REF/Aly. B)- The nuclear export of mRNAs generated from intronless genes is either due to recruitment of REF/Aly by CBP20, or by binding of SR proteins to the mRNA. In both cases, TAP/p15 facilitates nuclear export. CBC, Cap-Binding Complex.

junction complex (EJC) for intron-containing genes, by cleavage and polyadenylation at the 3' end, and by the binding of RNA-export adaptor proteins (reviewed in (63)). Each of these steps is carried out co-transcriptionally by distinct multi-protein complexes, which are physically and functionally coupled (64, 65). In a more recent model (Figure 3A), fully processed mRNPs generated from intron-containing genes are exported via 5' end recruitment (by CBP80) of the RNA export factor REF/Aly bound to UAP56 in the transcription elongation complex TREX which is stably associated with RNA probably by interacting with the EJC complex (55). REF/Aly would then interact with TAP/p15 that facilitates the export of mRNPs by interacting with nucleoporins via its C-terminal NES. For mRNPs generated from intronless genes, one pathway of export involves binding of SR proteins 9G8 and SRp20 to specific RNA sequences, which facilitate export by directly recruiting TAP/p15 (48, 66) (Figure 3B). In another alternative pathway REF/Aly would be recruited to intronless mRNAs by CBP20, which facilitates export by recruiting TAP/p15 (67) (Figure 3B). From the above models, it can be concluded that in mammalian cells, UAP56 and REF/Aly are essential mRNA export adaptors, while TAP/p15 is a promiscuous mRNA export receptor.

EB2, REF/Aly and TAP co-immunoprecipitate from living cells but this complex is disrupted by ribonuclease treatment, suggesting that it is stabilized by RNA (34). Interestingly, the REF/Aly interaction domain in EB2 was located in the leucine-rich region (Figure 2B) and

EB2-mediated mRNA export was severely impaired by deletion of this region. In *in vitro* binding assays, EB2 interacted with the C-terminal region of REF/Aly. The interaction was not disrupted by ribonuclease treatment, demonstrating a direct interaction between EB2 and REF/Aly *in vitro* (34). Thus EB2 appears to recruit TAP indirectly via REF/Aly to export mRNAs.

In a yeast two-hybrid screen, human OTT3 was identified as an EB2-interacting factor (68). OTT3/RBM15B is a member of the Spen family of proteins which include human OTT1/RBM15 and SHARP (69). These factors are characterized by several RNA Recognition Motifs (RRMs) located at their N-termini and by a C-terminal conserved SPOC (Spen Paralog Ortholog C-terminal) domain. Recently, RBM15 has been described as an mRNA export factor (51) that binds specifically via its RRM to the transport element RTE identified in mouse retrotransposon unspliced mRNAs and directly interacts with TAP/p15 via the SPOC domain, which facilitates the nuclear export of RTE-containing mRNAs. EB2 interacts with the SPOC domains of RBM15, OTT3 and SHARP but it is not known at present if OTT3 and SHARP are also RNA export factors. EB2 and TAP/p15 both interact with the SPOC domain of OTT1 but it is not yet known whether this is with the same or different regions in the SPOC domain. OTT3 interacts with EB2 via amino acids 10 to 40 at the N-terminal (Figure 2B), while RBM15 and SHARP interact with the C-terminus of EB2 (68). In conclusion, EB2 appears to export mRNAs to which it is bound, by directly contacting several RNA export adaptor proteins of the TAP/p15 pathway.

4.4 EB2 induces the cytoplasmic accumulation of mRNAs

In line with the observations that EB2 shares properties with mRNA export factors, EB2 has been shown to affect the cytoplasmic accumulation of both intronless and unspliced mRNAs. However, there is little agreement between the results obtained by different laboratories. One report suggested that EB2 was inducing the cytoplasmic accumulation of mRNAs generated from transfected intronless artificial reporter genes while inhibiting the export of mRNAs generated from transfected intron-containing artificial reporter genes (70). However, the same laboratory also showed that EB2 exported STAT1alpha and STAT1beta-spliced mRNAs in EBV-negative B cells (71). Other assays made use of a reporter gene called pDM128 (Figure 4A) (72), widely used to assess if a given protein has RNA export activity. This construction is an artificial gene with two exons and the HIV-1 *tat/rev* intron containing the CAT gene. The intron has suboptimal signals in the 3' splice site (73). Upon transient expression of pDM128 pre-mRNA in cells, the intron is excised prior to nuclear export of the RNA and little or no CAT protein is found in the cell extract. However, if RNA export factors are tethered to the intron via a specific RNA-binding element, then unspliced RNAs are exported and CAT protein can be detected in the cell extract (74). In this assay, EB2 greatly enhanced the export of pDM128 unspliced RNAs even in the absence of a specific RNA-binding motif (Figure 4A) (44). In another assay, the wild

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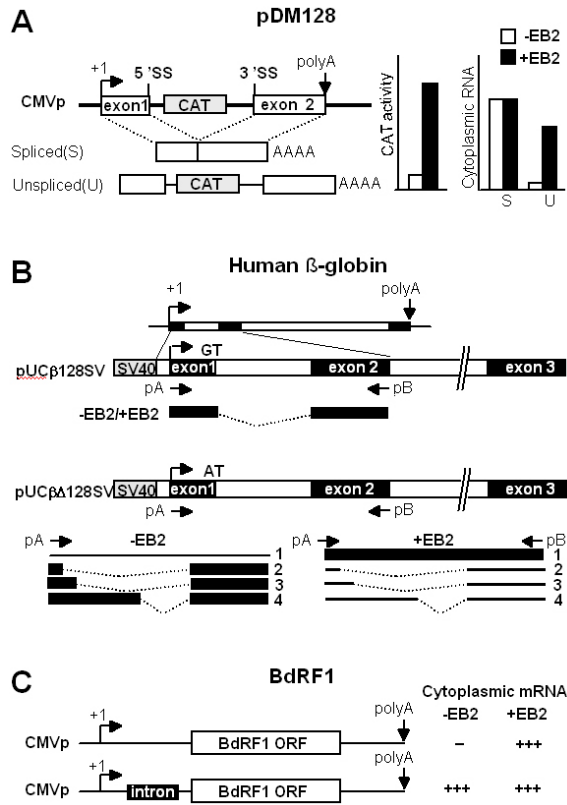


Figure 4. EB2 exports unspliced and intronless mRNAs. A)-Schematic representation of pDM128, of the CAT activity and of the spliced (S) or unspliced (U) cytoplasmic mRNAs expressed in the absence or presence of EB2. B)-Schematic representation of pUCbeta128SV and pUCbetaΔ128SV carrying the wt human beta globin gene or its beta thalassemia variant respectively, under the control of the SV40 early promoter. Only splicing of the first intron has been assessed by RT-PCR using primers pA and pB. Similar amounts of spliced wt beta globin mRNA is exported in the presence or absence of EB2 (-EB2/+EB2 under the reporter plasmid pUCbeta128SV). Four RNAs are expressed from the beta-thalassemia reporter plasmid, 1: unspliced RNA; 2,3 and 4: spliced RNAs generated by the use of three facultative cryptic 5' splice sites. In the presence of EB2, the unspliced RNA is the major RNA species exported (the thickness of the schematic mRNAs is directly proportional to the efficiency of their export). C)-The viral BdRF1 intronless reporter gene generates an mRNA that is only exported in the presence of EB2. No effect by EB2 on the export of the BdRF1 mRNA that has undergone splicing is detected.

type human beta-globin gene and its thalassemic allele were used (Figure 4B) (75, 76). The beta-thalassemic gene contains a G-to-A transition at position one of the first intron which causes the activation of three cryptic 5' splice sites. In this assay, EB2 had no effect on the nuclear export of wt spliced beta-globin mRNA, but strongly increased the export of unspliced beta-thalassemic RNA which was associated with a decreased export of spliced beta-

thalassemic mRNAs (Figure 4B) (44, 61). Most of the early and late EBV mRNAs are generated from intron-less genes, and in transient expression assays EB2 increases the export of some of these mRNAs (32, 77). However, when an intron was added to some of these otherwise intronless genes, then EB2 had no effect on the already efficient export of the mRNAs that had undergone splicing (Figure 4C) (32, 77). In conclusion, in transient expression assays, EB2 increases the cytoplasmic accumulation (export) of mRNAs generated from intronless genes as well as of unspliced mRNAs generated from introns containing genes but only if they contain facultative or cryptic splice sites.

It should be stressed that an EB2 region essential for its effect on the export of intron-containing and intronless mRNAs has been located in the 35 amino acids at the C-terminal (78). However, mutagenesis in this region mostly generated EB2 proteins that were abnormally distributed in the nucleus as dense separate foci, representing insoluble aggregates in which EB2 was inactive. Moreover, as compared to the diffuse nucleoplasmic localization of wt EB2, several overlapping deletions covering the 250 amino acids at the EB2 C-terminal generated the same phenotype - dense separate nuclear foci (Figure 5, our unpublished results) - and these EB2 mutants were inactive in an RNA-export assay (61). This suggests that the C-terminal part of EB2 is highly structured and that disruption of the structure induces aggregation and inactivation of the protein. It should also be noted that as EB2 has been reported to self-associate *in vitro*, via a region located in the 200 amino acids at the C-terminal (79), it is also possible that inactivation of self-association induces aggregation and inactivation of EB2 *in vivo*.

5. MOLECULAR GENETICS OF EB2

In order to study the function of EB2 in the context of the EBV productive cycle, an EBV mutant was generated in which the EB2 gene was deleted. The mutated viral genome was then introduced into 293 cells and one cell clone called 293_{BMLF1-KO} was selected. (80). In order to induce the productive cycle of EBV in cells latently infected with the virus, it is necessary and sufficient to transiently express the EBV transcription factor EB1/ZEBRA which is the activator of both EBV early gene expression, including that of EB2, and viral DNA replication. The amount of virions produced by the 293_{BMLF1-KO} cells can be evaluated by infecting B cells with the 293_{BMLF1-KO} supernatant. Due to the constitutive expression of the GFP protein from the EBV recombinant genome, infected cells are bright green under UV light and can be quantified by FACS analysis. In this assay, 293_{BMLF1-KO} cells harbouring the EB2-defective EBV genome only produced infectious EBV particles when both EB1 and EB2 were transiently expressed, demonstrating that EB2 is essential for the production of infectious virions (80).

In this transcomplementation assay, it was shown that most of the late viral mRNAs but only a subset of the early viral mRNAs are exported by EB2 (Figure 6). This was achieved by semi-quantitative RT-PCR analysis (77,

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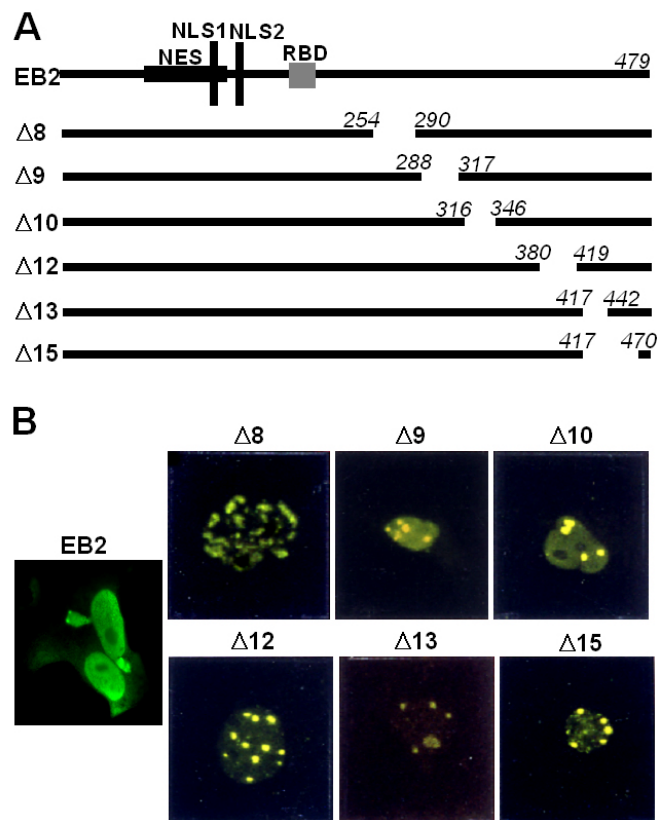


Figure 5. Deletions in the C-terminal half of EB2 induce its nuclear aggregation. A)- Schematic representation of EB2 C-terminal amino acid deletions. B)- Nuclear distribution in dense separate nuclear foci of EB2 mutated proteins expressed in Cos-1 cells, as visualized by indirect immunofluorescence.

80) and by oligonucleotide arrays (81). These results have important implications. Firstly, the inefficient export of the early mRNAs encoding the primase and the DNA polymerase in 293_{BMLF1-KO} cells expressing EB1 alone, was associated with poor (77) or absent (81) viral DNA replication. As viral DNA replication is a prerequisite for the expression of the late viral genes, the defect in late mRNA expression in the absence of EB2 could be the consequence of poor replication of the viral DNA. However, although transfection of primase and DNA polymerase expression vectors in 293_{BMLF1-KO} cells together with EB1 partially rescued DNA replication, the transfected cells produced no infectious virions (81). This could be due to the fact that the expression of the single stranded DNA-binding protein is also EB2-dependent (80). Thus expression vectors for the single stranded DNA-binding protein should have been co-transfected together with expression vectors for EB1, the primase, and the DNA polymerase, to efficiently rescue DNA replication. Nevertheless, about 12 late genes were detectably expressed in 293_{BMLF1-KO} cells in which DNA replication was partially rescued, indicating that replication *per se* increases the expression of some late genes and that the corresponding mRNAs were exported in the absence of EB2. It is noteworthy that their export was further increased by EB2. However, in conditions that partially rescued replication, the majority of the late mRNAs were

strictly dependent on EB2 to be exported. These results suggest that EB2 is required for the efficient export of the majority of the viral late mRNAs, and for the production of infectious virions. Secondly, expression in 293_{BMLF1-KO} cells of both EB1 and an EB2 mutant with the REF/Aly interacting domain deleted, induced viral DNA replication efficiently but very inefficiently rescued the production of infectious virions (80). This strongly suggests that EB2 mainly exports late viral mRNAs via the REF/TAP pathway and that the early mRNAs encoding proteins essential for the replication of viral DNA are exported by EB2 via another pathway. Thirdly, the absolute requirement of EB2 for the efficient production of infectious virions has also been linked to a defect in the intranuclear assembly of capsids but this was expected since EB2 is required for the efficient export of the mRNAs encoding the capsid proteins (77).

It is clear from the transcomplementation assay, that some early and most late EBV mRNAs are exported by EB2. Although most of these mRNAs are generated from intronless genes, EB2 also exports unspliced viral mRNAs. Indeed, the EBV BLLF1 late gene contains an intron and both spliced and unspliced BLLF1 mRNAs must be exported to generate respectively gp220 and gp340, two glycoproteins essential for the infection of cells by EBV (82). When 293_{BMLF1-KO} cells, were transfected with EB1

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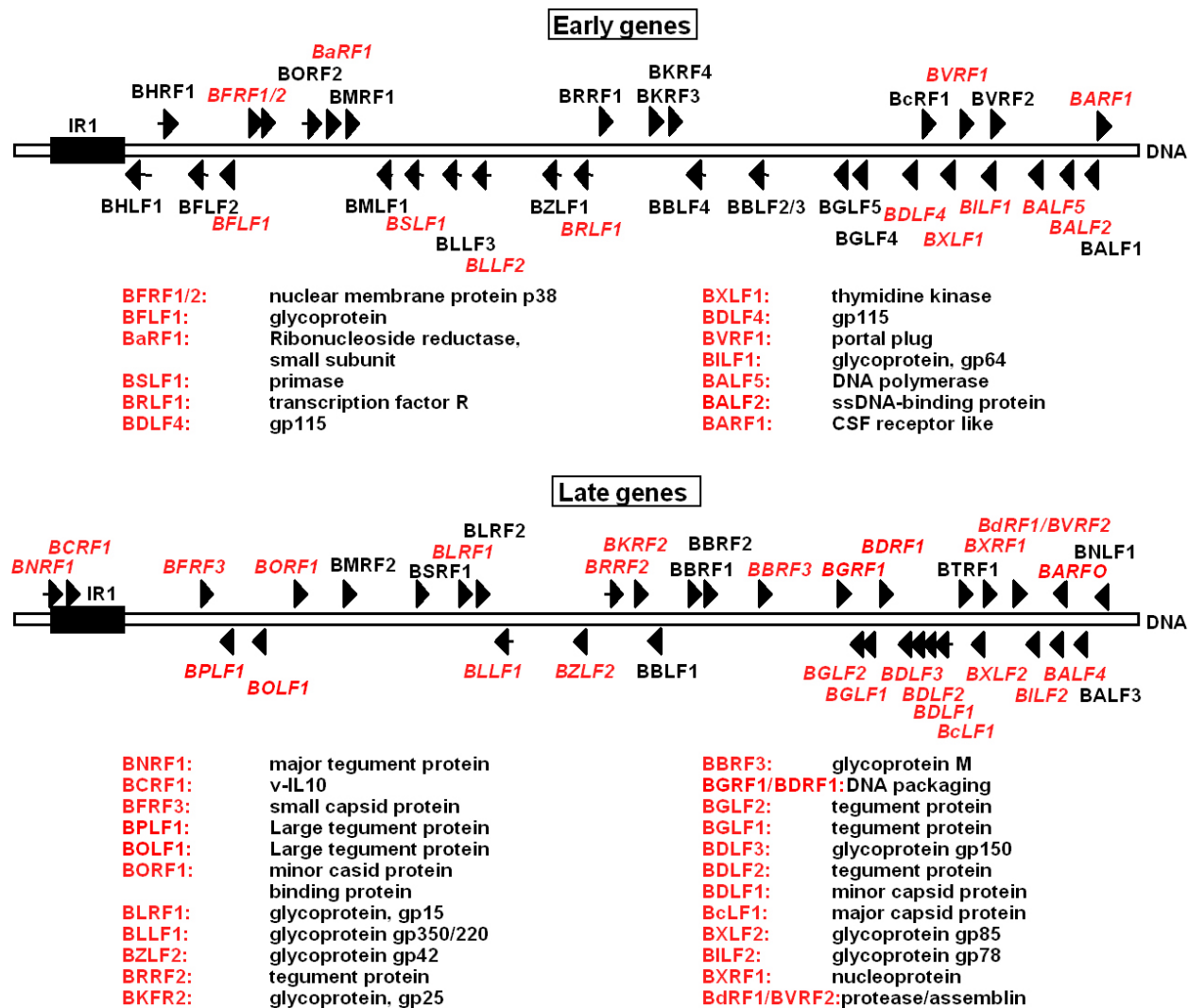


Figure 6. EB2 exports a subset of early and late viral mRNAs. The mRNAs exported by EB2 characterized by RT-PCR and oligonucleotide arrays, are indicated in red italics over and under the schematic representation of the viral genome (DNA). Their location and the orientation of their synthesis are indicated by black arrowheads. The protein encoded by each mRNAs exported by EB2 is also indicated. IR1: Internal Repeats 1.

alone, only the spliced mRNA was expressed and exported, whereas when both EB1 and EB2 were expressed the export of the unspliced mRNA was induced and the export of the spliced RNA remained as efficient as when EB1 was expressed alone (77). Therefore, EB2 also exports unspliced viral mRNAs carrying facultative splice sites, which is in agreement with what was observed in transient expression assays using heterologous genes (see 4.4).

Using the transcomplementation assay, it was also shown that the EB2 mutant lacking the RNA-binding domain did not rescue the production of infectious virions by 293_{BMLF1-KO} cells, suggesting that EB2-binding to RNA *in vivo* is required for the production of EBV-infectious virions (34). On the contrary, EB2 with the OTT3 N-terminal binding

domain deleted efficiently rescued the production of infectious viral particles (our unpublished results). However, since this EB2 mutant still potentially interacts with RBM15/OTT1, this result is not so surprising as these proteins could transcomplement each others function. Finally, none of the EB2 mutants generated in the C-terminal 35 amino acids that localized abnormally in the nucleus, rescued the production of infectious virions by 293_{BMLF1-KO} cells (78). In conclusion: 1) DNA replication and EB2 are both required for an efficient nuclear export of the majority of late viral genes, probably mostly mediated by the REF/TAP pathway; 2) EB2-mediated export of early genes probably depends on a pathway that does not involve REF; 3) many early and some late intronless viral mRNAs are efficiently exported in the absence of EB2 via pathways that have not as yet been identified.

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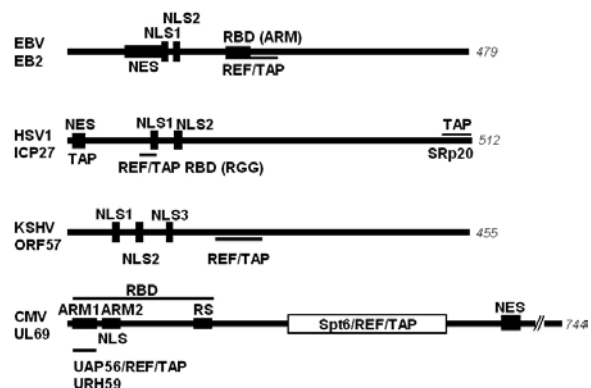


Figure 7. Functional domains and factors that interact with EB2 and with its functional homologues. Schematic representation of the functional domains of EB2, HSV-1 ICP27, KSHV ORF57 and CMV UL69 and of the regions that interact with cellular factors. RBD: RNA Binding Domain; ARM: Arginine Rich Motif; RGG: Arginine Glycine rich motif; RS: Arginine Serine rich motif.

6. HSV1 ICP27, KSHV ORF57 AND CMV UL69 DO NOT COMPLEMENT EB2 IN VIRUS PRODUCTION

Several human herpes viruses carry a gene in their genome whose product is functionally similar to EB2, for example, the Herpes Simplex virus type 1 (HSV1) protein ICP27 (47, 83, 84), the Cytomegalovirus (CMV) protein UL69 (85), and the Kaposi's Sarcoma-associated Herpes Virus (KSHV) protein ORF 57 (86) (Figure 7). At least for HSV1 and KSHV, the respective deletion of the ICP27 (87) and ORF57 (88) genes, abolished both the production of infectious viral particles and the cytoplasmic accumulation of specific viral mRNAs, demonstrating that, like EB2, ICP27 and ORF57 are essential viral factors.

Importantly, like EB2, HSV1 ICP27 (47, 84), CMV UL69 (89) and KSHV ORF57 (86) shuttle between the nucleus and the cytoplasm in a CRM1-independent manner. Like EB2, they also induce the cytoplasmic accumulation of some but not all viral mRNAs (85, 90, 91), by interacting with cellular adaptors of the TAP/p15 receptor pathway (Figure 7). The CMV UL69 protein interacts with the DEXD/H-box RNA helicase UAP56 (85), a cellular factor that in metazoans appears to interact with REF/Aly which in turn makes contact with TAP/p15. UL69 also interacts with the human homologue of yeast Spt6 (92), a histone H3-H4 chaperone that maintains native chromatin structure behind elongating RNA Pol II (93) and mediates nucleosome reassembly at promoters during transcriptional repression in yeast (94). Human Spt6 binds RNA Pol II as well as Iws1, which associate with REF/Aly (95). Thus interaction of UL69 with both Spt6 and UAP56 likely mediates the efficient recruitment of TAP/p15 for the export of intronless viral mRNAs. ICP27 interacts with REF and TAP/p15 (47, 84) and with the splicing factor SRp20 that binds directly to TAP, although a function of SRp20 in ICP27-mediated mRNA export has not been documented (96). The KSHV ORF57 also interacts with

REF and TAP/p15 (86). In addition, UL69 (97), ICP27 (83), EB2 (34) and ORF57 (86) bind to RNA *in vitro* and *in vivo*, but the binding is not sequence or structure specific. Surprisingly, RNA-binding of UL69 is not a prerequisite for the nuclear export of unspliced pDM128 mRNAs (97). Nevertheless, since EB2, ICP27, UL69 and ORF57 bind non-specifically to viral RNAs *in vivo* and recruit adaptor proteins to export RNAs via the REF/TAP pathway, it could have been expected that they would trancomplement each other in the export of viral mRNAs and in the production of infectious virions. Yet, when inserted into a HSV1 ICP27-null mutant, EB2 did not efficiently complement ICP27 in inducing the production of infectious virions (98). When provided in trans, neither ICP27, UL69 (80) nor KSHV ORF57 (88) complemented EB2, in inducing the production of infectious virions from 293_{BMLF1-KO} cells. These results suggest that some of the known RNA-binding proteins that coat cellular mRNPs during their biogenesis bind preferentially to specific sub-populations of viral mRNPs and specifically recruit one or the other of the herpes virus export factors. The observation that EB2, ICP27, UL69 and ORF57 are absolutely required for the export of viral mRNAs and the production of infectious virions, strongly suggests that viral mRNAs inefficiently bind REF or other RNA export adaptors in the absence of viral export factors.

7. THE mRNA EXPORT FUNCTION OF EB2 IS REGULATED BY PROTEIN KINASE CK2

EB2 is a nuclear phosphoprotein (99) which is phosphorylated by protein kinase CK2 (24) but whether CK2 binds to EB2 or whether EB2 phosphorylation by CK2 affects its mRNA export function had not been established. Protein Kinase CK2 is a heterotetrameric complex consisting of a stable association of two catalytic subunits (alpha or alpha') and two regulatory beta subunits (Figure 8), that phosphorylates many various substrates localized in different cellular compartments (for a review see, (100)). However, free populations of CK2 subunits can also exist within the same cell (101). Both CK2 subunits are independently targeted to the nucleus where they exhibit differential mobility and CK2alpha but not CK2beta, shuttles between the nucleus and the cytoplasm. Thus, CK2 can act not only as a hetero-tetrameric holoenzyme but also as free populations of both subunits, either alone or in association with different partners, and in different subcellular compartments.

Our recent *in vitro* studies (102) show that both the CK2 alpha catalytic subunit and the beta regulatory subunit bind individually but more efficiently as the alpha2beta2 heterotetrameric holoenzyme, to the region containing the N-terminal NES of EB2 (Figure 8). It is evident that docking of CK2 or of CK2 subunits in the region containing the NES does not interfere with the nucleo-cytoplasmic shuttling of EB2. It is also unlikely that CK2 alpha participates in the nucleo-cytoplasmic shuttling of EB2 since EB2 shuttling is resistant to the CRM1 inhibitor leptomycin B (34) whereas this inhibits CK2 alpha shuttling (101). The alpha subunit and more efficiently the beta subunit also bind individually *in vitro* at

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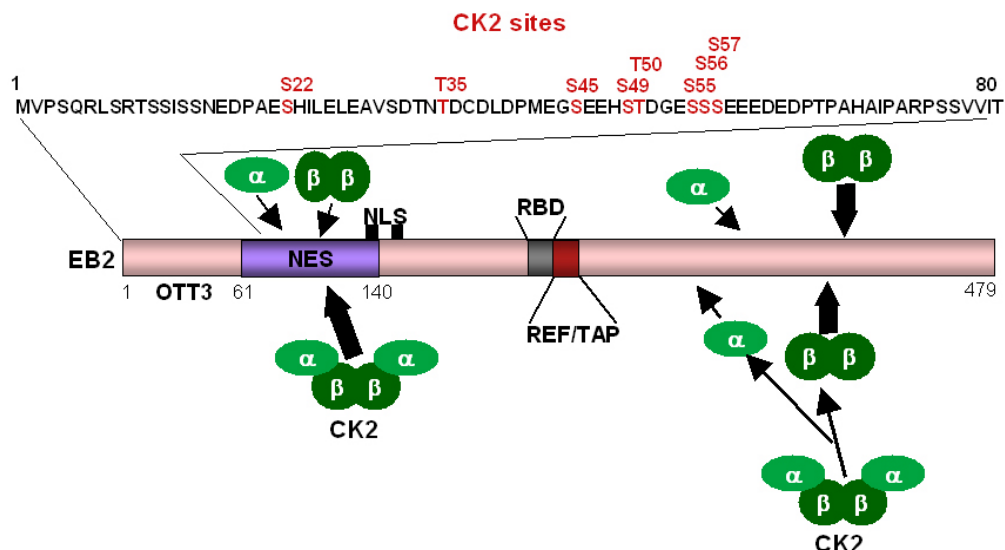


Figure 8. EB2 interacts with and is phosphorylated by protein kinase CK2. Individual CK2 subunits interact with both the N- and C-terminus of EB2 *in vitro*. The heterotetrameric CK2 also binds to the N-terminus of EB2, whereas incubation of the heterotetrameric CK2 with the C-terminus of EB2 leads to efficient binding of the beta-subunit and weak binding of the alpha subunit (the thickness of the arrows is directly proportional to the efficiency of binding). The N-terminal CK2 phosphorylation sites are shown, including serines 55, 56 and 57. The EB2 regions interacting with OTT3, REF and TAP are also shown.

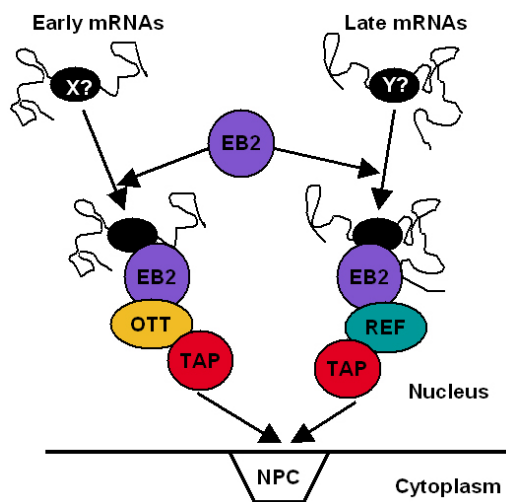


Figure 9. Models for EB2-mediated mRNA export. The viral mRNAs do not bind REF efficiently or other export adaptors in the absence of EB2. EB2 is recruited to early and late intronless mRNAs by different factors that by as yet unknown mechanisms, direct the binding of EB2 to RNAs, the recruitment by EB2 of different export adaptors and the export of the mRNAs.

the C-terminus of EB2 (Figure 8). Interestingly, when the holoenzyme is incubated *in vitro* with the EB2 C-terminal 295 amino acids, the beta subunit binds much more efficiently than the alpha subunit. This suggests that interaction of CK2 with the C-terminus of EB2 destabilizes the heterotetrameric enzyme, leading to efficient binding of the beta subunit alone, and weak binding of the alpha subunit, alone and/or as a heterotetrameric complex.

Only the N-terminus of EB2 is phosphorylated by CK2 *in vitro*. All the phosphorylation sites are located

between amino acids 1 and 60, upstream of the CK2 binding site, and include the three serines S55, S56 and S57 located next to the NES (Figure 8). EB2 and CK2 co-purify as a complex from living cells, and in this complex, CK2 phosphorylates EB2 at all N-terminal sites. However, among several phosphorylation mutants generated, only the substitution of the three serines 55, 56 and 57 by alanines (mutant EB2S3A) had a functional effect unlike their replacement by glutamates (mutant EB2S3E). Indeed, the EB2 phosphorylation ablation mutant EB2S3A inefficiently rescued the production of infectious virions by 293_{BMLF1}KO

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cells, whereas the EB2 phosphomimetic mutant EB2S3E was as efficient as wild type EB2. Although the three serines 55, 56 and 57 are located next to the NES, their mutation to alanine or glutamate residues did not affect the nucleo-cytoplasmic shuttling of EB2. Rather, and in line with the mRNA export function of EB2, the incapacity of EB2S3A to rescue infectious virus production was associated with the inefficient cytoplasmic accumulation of several late mRNAs, including those encoding the EBV capsid proteins. This strongly suggests that the mRNA export function of EB2, and as a consequence the production of infectious virus, is regulated by protein kinase CK2. However, why CK2 phosphorylation of EB2 at the level of at least one of the Ser-55, -56 and -57 residues, is critical for the efficient export of several viral late mRNAs is not known at present. It is noteworthy that phosphorylation *per se* is not required, since EB2S3E is functionally equivalent to EB2wt. It is also not known what function is associated with the binding of the alpha and beta subunits to the EB2 C-terminus.

8. EB2 INTERACTS WITH THE PML BODY PROTEIN Sp110b

The nucleus of mammalian cells contains several nuclear domains or foci, which contain multiple proteins. These nuclear bodies include nucleoli, PML bodies (also called ND10), splicing speckles, Cajal bodies and paraspeckles. The proteins found in these subnuclear structures interact dynamically with one or the other of these bodies, and are involved in different nuclear functions, such as ribosome biogenesis, transcription and RNA splicing. EB2 has been found to interact with Sp110b, a splice variant of the PML body Sp110 component (103). PML bodies contain about 80 proteins involved in transcription, DNA repair, splicing, histone modification, and chromatin remodeling. These proteins interact dynamically with the PML bodies and can also be found in the nucleolus, in splicing speckles and at telomeres. More importantly, PML bodies have been described as facilitating the transcription and replication of double-stranded DNA viral genomes (reviewed in (104)).

EB2/SM has been found to interact with and to induce the expression of Sp110b (105) which has been shown to act as a transcriptional repressor of retinoic acid receptor alpha (106). The interaction between EB2/SM and Sp110b was first observed in a yeast two-hybrid screen and confirmed by co-immunoprecipitation from transfected HeLa cells. In transient expression assays, it was shown that Sp110b had no effect on the expression of intronless heterologous reporter genes, but enhanced EB2/SM-mediated activation of the target genes. The effect was post-transcriptional, and Sp110b prolonged the half-life of a target intronless heterologous mRNA in the presence of EB2/SM. It was further shown by transient expression assays, that EB2/SM induced the expression of Sp110b in both B lymphocytes and epithelial cells, and that Sp110b was induced during EBV replication in Burkitt's lymphoma cells and gastric epithelial cells. Finally, siRNA partial inhibition of Sp110b expression modestly but clearly decreased the expression of one EBV gene, the DNA

polymerase processivity factor BMRF1, in EBV-infected B cells in which the lytic cycle was induced by TPA + butyrate. Taken together, these results suggest that Sp110b stabilizes some EBV mRNAs by interacting with EB2, leading to enhanced EB2-mediated export of these mRNAs in EBV infected cells undergoing lytic replication. However, apart from the BMRF1 mRNA, the Sp110b/EB2 viral mRNA targets have not as yet been defined.

9. PERSPECTIVES

Although it is now well documented that EB2 has the properties of an RNA export factor, the final proof that EB2 is an RNA export factor would be to inject EB2 into the nucleus of *Xenopus laevis* oocytes, together with ³²P-labeled mRNAs which are known to be exported or not by EB2, in the presence or absence of an excess of the TAP binding RNA motif CTE. In this assay, would EB2 specifically export mRNAs? Several other important questions remain unanswered. There appear to be several NESs in EB2. Where are they precisely located and what are the cellular factors facilitating EB2's shuttling? EB2 binds RNA *in vitro* and *in vivo*, but no specific sequence or structure required for binding have been identified. What makes an RNA a target for EB2-mediated mRNA export? Is EB2 recruited co-transcriptionally to the viral mRNPs and if so by which complex? EB2 interacts with several cellular RNA export factors of the TAP/p15 pathway, REF/Aly, OTT1 and perhaps OTT3 (Figure 9). What is the contribution of these factors to the EB2-mediated nuclear export of viral mRNPs? In particular, which EBV mRNAs are exported in 293_{BMLF1-KO} cells expressing EB1 and EB2 with the regions interacting with REF/TAP, OTT3 and/or OTT1 deleted? The three-dimensional structure of EB2 or EB2 sub-domains should be established alone or in association with RNA or with the SPOC domain of OTT3 and OTT1. How does the phosphorylation of EB2 on serines 55, 56 and 57 control its function and how is this phosphorylation itself regulated? Is docking of CK2 essential for EB2 phosphorylation? Does EB2-bound CK2 phosphorylate viral and/or cellular proteins? Last but not least, why is EB2 essential for the nuclear export of just some viral mRNPs, and why can't its function be transcomplemented by HSV-1 ICP27 or CMV UL69? These are challenging studies for the future.

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Abbreviations: ARM: Arginine Rich Motif, AP-1: Activating Protein 1, CAT: Chloramphenicol Acetyl Transferase, CRM1: Chromosomal Maintenance Region 1, EJC: Exon Junction Complex, GST: Glutathione-S-transferase, NPC: Nuclear Pore Complex, RBD: RNA-Binding Domain, RRM: RNA Recognition Motif, RXP: Arginine-X-Proline

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