

# VCR X86 X64 13102011 20 ^HOT^

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## VCR X86 X64 13102011 20

RedGiant.Net REX Win32 Codec: VCR x86 x64 13102011 20 SVCDecoder.Net Win32 VCR x86 x64 13.10.2011 20 ComputerVirus.Win For dummies 2015...Mutations of the phosphoprotein associated with retention in the endoplasmic reticulum of the hepatitis B virus E2 protein affect its endoplasmic reticulum retention. A mutation of the phosphoprotein associated with retention in the endoplasmic reticulum (pEER) (38K) domain of the hepatitis B virus (HBV) E2 protein from an arginine residue (in the corresponding residue of the E1 protein) to a serine residue (pEER-R96S) also resulted in an increase in the endoplasmic reticulum retention of the mutant protein. Mutations of the pEER domain from an arginine residue to a serine residue (pEER-R96S) also result in an increase in endoplasmic reticulum retention of the E2 protein. The increased endoplasmic reticulum retention of the pEER-R96S protein was not due to slower turnover in the endoplasmic reticulum because the degradation rate of the pEER-R96S protein in the endoplasmic reticulum was not increased. By comparing the accumulation of the mutant and wild-type E2 proteins during transient expression in HepG2 cells, we found that the accumulation rate of the mutant protein in the endoplasmic reticulum was reduced by about 50%. Mutations of the pEER domain of the E2 protein from an arginine residue to a serine residue (pEER-R96S) also results in an increase in endoplasmic reticulum retention of the E2 protein. To determine whether the decrease in the accumulation rate of the mutant protein could be attributed to defects in its export to the plasma membrane or to decreased stability in the endoplasmic reticulum, we compared the export and degradation rate of the mutant and wild-type E2 proteins after they had reached their steady-state level of accumulation in the endoplasmic reticulum. Although the mutant and wild-type E2 proteins were equally stable in the endoplasmic reticulum, the mutant protein was not exported to the plasma membrane as efficiently as 04aeff104c

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