Figure S1 (related to figures 1 and 3)



(A) THP-1 cells stably expressing the indicated or control shRNA were differentiated with PMA for 48h and infected with HIV-1Luc. After 2 days, cells were lysed for luciferase luminescence. Average of two independent experiments are shown. (B) Western blot showing SAMHD1 knockdown by shRNA in differentiated THP-1 cells. (C) Differentiated THP-1 cells were infected with HIV-1Luc for 2 h. Cells were washed vigorously three times, lysed and p24 ELISA performed on the cell lysates.

Figure S2 (related to figure 2)



(A) Cyclin L2 and DCAF1 show partial colocalization in the nucleus. HeLa cells were transfected with myc-Cyclin L2 and stained for myc and endogenous DCAF1. Blue staining in the top panel represents DAPI (B) THP-1 cells were lysed and immunoprecipitation performed with Cyclin L2 antibody and Western blots probed for Cyclin L2, DCAF1 and DDB1.

Figure S3 (related to figure 3)

Cyclin L2 does not affect HIV replication in HeLa and 293T cells



(A) 293T cells were transfected with control or siRNA to Cyclin L2 for 48h and WB performed. (B) 293T cell were transfected with control or Cyclin L2 siRNA. After 48h, cells were infected with HIV-1luc and viral replication determined by luciferase luminescence in cellular lysates (C) 293T cells were transfected with control or Cyclin L2 plasmid and Western blot performed after 48h. Antibodies to epitope tag (myc) and Cyclin L2 on the same blot are shown. (D) 293T cells transfected with Cyclin L2 or control plasmid for 48h were infected with HIV-1luc and viral replication measured by luciferase in cellular lysates (E) HeLa cells were transfected with control or siRNA to Cyclin L2 for 48h and WB performed (F) HeLa cells were transfected with control or Cyclin L2 siRNA. After 48h cells were infected with HIV-1luc and viral replication determined by luciferase luminescence in cellular lysates. Data indicate means; error bars indicate \pm SEM (n \ge 3).

Figure S4 (related to figure 4)



Cyclin L2 shows significant nuclear colocalization with SAMHD1. HeLa cells were transfected with SAMHD1 overnight and immunostained for SAMHD1 and endogenous Cyclin L2.

Figure S5 (related to Figure 5)



(A) 293T cells were transfected with increasing concentrations of SAMHD1 and 1ug of Cyclin L2 for 48h and the expression levels of the two proteins determined by Western blot. (B) Stable THP-1 cells expressing control or Cyclin L2 shRNA were differentiated for 48h in PMA and WB performed on a Phostag gel for SAMHD1 levels.