Proviral Structure, Chromosomal Location, and Expression of HERV-K-T47D, a Novel Human Endogenous Retrovirus Derived from T47D Particles

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We previously described that type B retrovirus-like particles released from the human mammary carcinoma cell line T47D are pseudotypes and package retroviral RNA of different origins (W. Seifarth, H. Skladny, F. Krieg-Schneider, A. Reichert, R. Hehlmann, and C. Leib-Mösch, J. Virol. 69:6408–6416, 1995). One preferentially packaged retroviral sequence, ERV-MLN, has now been used to isolate the corresponding full-length provirus from a human genomic library. The 9,315-bp proviral genome comprises a complete retroviral structure except for a 3′ long terminal repeat (LTR) truncation. A lysine tRNA primer-binding site and phylogenetic analyses assign this human endogenous retroviral element, now called HERV-K-T47D, to the HML-4 subgroup of the HERV-K superfamily. The gag, pol, and env genes exhibit 40 to 60% amino acid identity to HERV-K10. HERV-K-T47D is located on human chromosome 10, with five closely related elements on chromosomes 8, 9, 15, 16, and 19 and several hundred HERV-K-T47D-related solitary LTRs dispersed over the human genome. HERV-K-T47D-related sequences are detected in the genomes of higher primates and Old World monkeys but not in those of New World monkeys. High HERV-K-T47D transcription levels were observed in human placenta tissue, whereas transcription in T47D cells was strictly steroid dependent.

Human endogenous retroviral sequences (HERVs) are inherited genomic elements with structural features of integrated retroviruses. To date, HERVs are estimated to comprise at least 1% of the human genome (for reviews, see references 14, 16, and 44). The biologically most active HERVs are members of the HERV-K superfamily (for a review, see reference 20). Members of this family are characterized by the presence of primer binding sites (PBSs) for lysine tRNA, hence the designation K. They represent about 70 to 100 elements and a large number of solitary long terminal repeats (LTRs) in the human genome. These elements are related to type A, B, and D retroviruses and have been classified by alignments of short stretches of the reverse transcriptase (RT) domain into six different groups (HML-1 to -6) (23). The members within a subgroup are more than 85% identical, whereas the intersubgroup similarity does not exceed 75%. To date, full-length proviral elements from only subgroups HML-2 and -6 have been isolated and completely sequenced.

The group HERV-K(HML-6), which is the least closely related to mouse mammary tumor virus, comprises about 30 to 40 members with 40 to 68% nucleotide sequence similarity to mouse mammary tumor virus and intracisternal type A particles of the mouse and hamster (24). In addition to proviral sequences, about 50 solitary HML-6-related LTRs are found per haploid genome.

The HERV-K(HML-2) group consists of approximately 30 members with full-length genomes, a few elements with large deletions (23, 25, 31), and an estimated 10,000 to 25,000 solitary LTRs distributed throughout the human genome (15). The prototype HERV-K(HML-2) provirus is HERV-K10, which to date is the only completely sequenced full-length provirus of this group (31). HERV-K10 and most HERV-K10-related proviruses harbor a characteristic deletion of 292 nucleotides (nt) leading to a defective genome with a polymerase gene fused to the envelope gene. However, transcripts of other HERV-K-related proviruses with uninterrupted pol and env open reading frames have been detected in human teratocarcinoma cell lines (18, 19, 42), and gag and pol gene products of HERV-K(HML-2) family members have been demonstrated to be enzymatically active (12, 28, 37). The env gene of HERV-K-IDDM, which was isolated from patients with acute-onset type I diabetes, was found to encode an endogenous superantigen (4). These studies suggest that some proviruses of the HERV-K superfamily have the potential to encode functional retroviral enzymes, possibly even with sufficient genetic information for the formation of retrovirus-like particles, which have been observed in normal human placenta, oocytes, and fetuses (9, 13, 21, 26), in both malignant and nonmalignant breast tissue samples (1, 10, 11, 27), and in germ cell tumors or cell lines derived from these tissues (17). However, there remain many questions with regard to the biological significance or function of these particles, particularly since they appear to be generated by complementation between several expressed HERVs, resulting in pseudotype particles with retroviral RNA of different types (2, 32, 38). Such packaging mechanisms could lead to unforeseen consequences in the use of retroviral vectors in gene therapy or following interspecies organ transplants.

Previously, using degenerate primers from a conserved region of retroviral pol genes (39), we repeatedly amplified three different retroviral sequences from particles released by the
human mammary carcinoma-derived cell line T47D (38). One predominant sequence showed about 65% sequence identity to HERV-K10 within the RT region. By screening a human genomic library with the amplified product, we isolated a proviral pol sequence which we preliminarily termed ERV-MLN. The question was whether ERV-MLN is derived from an endogenous provirus with functional retroviral gene products, particularly with the packaging capabilities of Gag proteins. Therefore, we completely analyzed its proviral structure and genomic organization. Sequence comparisons assigned this novel HERV to the HML-4 subgroup of HERV-K elements. We also determined the chromosomal location and expression pattern of the provirus, now called HERV-K-T47D.

**Classification of the HERV-K-T47D provirus.** We previously isolated and cloned a human endogenous retroviral RT-related sequence from particles released by the human breast cancer cell line T47D. This pol fragment was used to isolate from a human genomic library, as described previously, a number of hybridizing λ clones (38) that were entirely sequenced by the dideoxy chain termination method (36). Two overlapping λ clones now revealed that this element, previously termed ERV-MLN, comprises an almost full-length proviral structure with an overall length of 9,315 bp (Fig. 1). Next to the 5' LTR (nt 1 to 943) is a putative tRNA PBS (nt 946 to 963) which, despite a 3-bp mismatch, is most closely related to the complementary sequence of the 3' end of human lysine tRNA (CUU anticodon) (Fig. 2A). The putative PBS is identical to that found in HERV-KC4 (6) and is closely related to the PBSS of other HERV elements belonging to the HERV-K superfamily. Therefore, ERV-MLN, a human endogenous retrovirus with lysine tRNA as the most likely primer for reverse transcription originating from retrovirus-like particles released by the T47D cell line, is now referred to as HERV-K-T47D.

To further classify HERV-K-T47D within subgroups HML-1 to -6, so far characterized by a 242-bp stretch of the RT domain (23), we aligned the subgroup sequences with the corresponding HERV-K-T47D region by using the software package Gene Works (IntelliGenetics, Inc.). Sequence comparison revealed that HML-4.1 is the most closely related sequence, showing about 80% nucleotide homology. Therefore, HERV-K-T47D is the first identified full-length prototypic element of the HERV-K(HML-4) subgroup.

**Genomic organization and coding regions.** Since a full-length HML-4 group provirus has not yet been identified, we used the well-characterized HERV-K10 (31), a member of the HML-2 subgroup, for alignments with a computer-assisted translation of the complete nucleotide sequence of HERV-K-T47D in order to analyze its genomic organization and identify putative reading frames. At the amino acid level, this revealed an HERV-K-T47D retroviral structure, comprising gag, pr, pol, and env genes (Fig. 2A), with six regions showing significant protein similarity to HERV-K10, ranging from 40 to 60% homology (boxes A to F in Fig. 1 and 2A). Within the HERV-K-T47D gag gene, which tentatively extends from nt 1095, based on comparison with HERV-K10 and the location of a suitable methionine, to nt 3480, two regions with significant amino acid similarity to HERV-K10 were found (box A, nt 1058 to 1385, 40% homology; and box B, nt 2416 to 3308, 40% homology). As shown in Fig. 2A, box A corresponds to the amino-terminal part of the matrix protein, whereas box B comprises the carboxy-terminal half of the capsid (CA) protein and almost the entire nucleocapsid (NC) protein. These regions are separated by 1,235 nt displaying no significant nucleotide (or, hence, amino acid) identity to the corresponding region of HERV-K10, which is shorter and comprises only 627 nt. FASTA database searches based on the differing 608 nt of this sequence revealed an 89% nucleotide homology in a 203-bp overlap with a human CpG island (3). Within the NC protein of HERV-K-T47D, two Zn finger domains of the CV4CV4H4CV4 type were identified (nt 3114 to 3155 and nt 3237 to 3275). The first motif is defective, lacking the initial Cys, whereas the second Zn finger is intact. A third conserved HERV-K-T47D region (box C in Fig. 1 and 2A) extends from nt 3510 to 4250. It exhibits 59% amino acid identity to the corresponding region of HERV-K10 and comprises the complete retroviral dUTPase (nt 3510 to 3883) and part of the retroviral protease (nt 3884 to 4340).

Retroviral pol genes are generally the most conserved sequences among retroviruses (22). This concurs with our observation that a 2.5-kb stretch of HERV-K-T47D pol (nt 4320 to 6998) shows a 60% overall amino acid identity to HERV-K10 (Fig. 2A, boxes D and E). The RT (nt 4341 to 5184) exhibits 65% identity, including some short stretches with almost absolute identity, while the tether region (nt 5185 to 5672), which connects the RT and RNase H protein domains, is less conserved (47% amino acid identity). RNase H (nt 5673 to 6086) shows 50% identity to HERV-K10, including a common feature of retroviral RNase H proteins, the DEDD motif.

The env gene of HERV-K-T47D shows the least homology to HERV-K10, with the exception of the transmembrane domain (TM). At the amino terminus of the HERV-K-T47D TM, a region with 59% amino acid identity to HERV-K10 is found (Fig. 1 and 2A, box F, nt 8347 to 8991). Specifically, two clusters of hydrophobic amino acids (nt 8401 to 8478 and 8931 to 9068)
FIG. 2. (A) Nucleotide sequence of HERV-K-T47D proviral DNA. LTRs are enclosed in brackets, and the inverted termini TGT and ACA are indicated by arrows. Transcriptional regulatory sequences, i.e., c/EBP, Gfi-1, AP-1, Ik-1, a glucocorticoid-responsive element (GRE), enhancer-like elements, a putative TATA box, a polyadenylation signal, and polyadenylation sites (CA and TA), are underlined once and labeled above. PBS and the polypurine tract (ppt) are double underlined. Sequence complementary to the 3' end of human lysine tRNA is depicted below the PBS sequence, with lowercase letters being used for mismatches. Translated amino acid sequences with significant homology to HERV-K10 (31), shown under the nucleotide sequence in the six shaded boxes A to F, are those of \( \text{gag} \) (box A, 40% identity; box B, 49% identity), dUTPase-protease (box C, 59% identity), RT-RNase H (box D, 59% identity), integrase (box E, 59% identity), and \( \text{env} \) (box F, 58% identity). Frameshifts in the amino acid sequence are indicated with slashes; asterisks correspond to stop codons.

Conserved zinc finger motifs (type \( CX_2CX_4HX_4C \)) in the NC (box B) region are marked by underlining of the corresponding amino acids. (B) Alignment of putative regulatory elements of the HERV-K-T47D 5' LTR with corresponding elements from solitary HERV-K-T47D-related LTRs of higher primates (33). Asterisks indicate binding sites which would not have been found with the default parameters of MathInspector (34). However, they were found when a lower threshold was used. Dots and dashes show identical and missing nucleotides, respectively. Under the binding site designations are search string variables used by the program MathInspector. IR, inverted repeat.
to 8970) are highly conserved (86% identity). The 3′ end of HERV-K-T47D env is followed by a polyurine tract (nt 9046 to 9059), which is a conserved motif of the retroviral env-LTR border. Despite their well-defined structure, the coding regions of HERV-K-T47D are interrupted by nonsense and frameshift mutations.

The putative LTRs of HERV-K-T47D were defined by aligning the sequences flanking the proviral coding regions at the 5′ and 3′ ends. Sequence repeats of 254 bp which differ from one another in 23 positions (91% homology) were identified. However, database searches revealed that the 943-bp region from the 5′ end of HERV-K-T47D exhibits 70% homology to a solitary retroviral LTR sequence at the human RNU2 locus on chromosome 17q21 (33). This LTR can be traced back to a complete retroviral element of 6 kb which still exists in the corresponding chromosomal locus of the baboon.

During primate evolution, excision of the provirus by homologous recombination created the solitary LTR now found in the genomes of the chimpanzee, gorilla, orangutan, and human (33). This LTR is considered to be associated with the concerted evolution of the tandem array encoding U2 snRNA. Direct sequence alignment of this solitary RNU2 LTR with the ends of HERV-K-T47D revealed that its 5′ LTR is intact whereas the 3′ LTR is truncated after nt 254. The 5′ LTR is bordered by short inverted repeats (TGT…ACA) and is followed by an untranslated leader sequence of 150 bp (nt 944 to 1094) containing the PBS (nt 946 to 963). Several potential regulatory elements were identified by using the program ModellInspector (8). Putative binding sites for transcription factors C/EBP, G6-1, AP1, and Ik-1 were detected within the U3 region (Fig. 2A and B) by using the program MathInspector (34). These sequences were found to be conserved in solitary human and various solitary primate RNU2 LTRs (Fig. 2B). However, a putative TATA box at position 462 of HERV-K-T47D is not present in those solitary LTRs. A glucocorticoid-responsive element and two enhancer-like structures were also tentatively assigned (29). A polyadenylation signal [poly(A)] was detected in the 5′ LTR but not in the 3′ LTR, which is truncated in this region. Therefore, a poly(A) signal located either within the coding region of the provirus or within 3′ cellular flanking sequences may be used to generate HERV-K-T47D mRNA. To examine these possibilities, we screened a cDNA library from steroid-induced T47D cells with an HERV-K-T47D LTR probe (Fig. 1, probe 1) generated by PCR with forward primer CCGAGGCAGAGACTGAGGC CAC (nt 25 to 47) and reverse primer ACTTCTCACAATGT CCCTTCAGC (nt 232 to 254). We were not able to isolate an HERV-K-T47D cDNA by this method, but we obtained several clones containing cellular sequences which are polyadenylated by solitary HERV-K-T47D-related LTRs (data not shown). Based on these clones, we identified two possible poly(A) addition sites (CA and TA) within the HERV-K-T47D LTR. The poly(A) addition site observed in the majority of cDNA clones was used to assign the R-US5 border (Fig. 2A).

Chromosomal location and evolution of HERV-K-T47D and related elements. Southern blot analysis was performed under high-stringency conditions as described previously (38) to determine the copy number and chromosomal location of HERV-K-T47D and closely related sequences in the human genome. HindIII-digested DNA from a panel of 24 human-rodent monochromosomal hybrid cell lines (mappan panel no. 2; NIGMS Human Genetic Cell Repository, Camden, N.J.) was hybridized to an HERV-K-T47D pol DNA fragment (Fig. 1, probe 2). The observed banding pattern suggests that HERV-K-T47D is located on human chromosome 10. Furthermore, five related elements, probably representing other members of the HERV-K(HML-4) family, could be assigned to chromosomes 8, 9, 15, 16, and 19. Southern blot analysis of human DNA samples digested with a set of restriction enzymes revealed that in addition to those proviral sequences, several hundred solitary HERV-K-T47D LTRs may exist in the human genome (Fig. 3A). As is known from studies of HERV-K(HML-2) (15) and HERV-H elements (7), multiple solitary LTRs are a common feature of HERV families. As evolutionary relics, they reflect high-level retrotransposon activity and subsequent homologous recombination during evolution.

To investigate HERV-K-T47D evolution, we analyzed DNAs of Old and New World monkeys and higher primates by performing Southern blot hybridization under relaxed hybridization conditions as described previously (38). High-molecular-weight DNA was digested with HindIII and probed with the 2.9-kb HERV-K-T47D pol fragment (Fig. 1, probe 2). A strong signal of the same size (2.9 kb) was detected in restricted DNA from both the human and the orangutan (Fig. 3B, lanes 1 and 3). DNA derived from the chimpanzee resulted in two smaller bands (1.3 and 1.4 kb) of similar intensity (lane 2), suggesting the presence of an additional HindIII restriction site in this element. In Old World monkeys (lanes 4 to 6), a series of weak signals differing in size (approximately 2.0, 2.4, and 3.5 kb) were detected, while DNA from the New World monkey genus Aotes (lane 7) gave no detectable hybridization signal. These data concur with previous findings indicating that most HERV elements arose early in primate evolution (for a review, see reference 16).

Transcription of HERV-K-T47D in human tissues. Regardless of whether active, functional proteins are encoded, a crucial role of HERVs may be their ability to act as promoters of either immunologically related retroviral antigens or cellular genes or, conversely, to act as premature transcription terminators. Since the production and release of T47D particles is steroid dependent (11, 32, 38), T47D cells were treated with 10⁻⁹ M estrogen for 48 h, at which time was added 10⁻⁸ M progesterone, with subsequent incubation for 24 h (11, 30).
Total RNA was prepared from steroid-treated and untreated cells in accordance with a CsCl ultracentrifugation protocol (35), separated by denaturing 1% formamide–agarose gel electrophoresis, transferred to Zeta-Probe membranes (Bio-Rad, Munich, Germany) by the vacu blot procedure (VacuGene XL; Pharmacia/LKB, Freiburg, Germany), and hybridized with a 

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\text{[32P]-labeled HERV-K-T47D fragment (Fig. 1, probe 2)}
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under high-stringency conditions (5 \( \times \) SSC [1 \( \times \) SSC is 0.15 M NaCl plus 0.015 M sodium citrate], 1% sodium dodecyl sulfate, 5 \( \times \) Denhardt’s solution, and 100 \( \mu \)g of denatured sheared herring sperm DNA per ml for 16 h at 65°C). Expression of HERV-K-T47D was found exclusively in full-term placental tissue (Fig. 4B). The high-stringency conditions revealed a high level of transcription exclusively in full-term placental tissue (Fig. 4B). The signal corresponds in size to the band obtained from T47D RNA after steroid treatment (Fig. 4A). With the exception of the breast carcinoma cell line T47D, a correlation between HERV-K-T47D expression and the occurrence or progression of malignancy was not found. No HERV-K-T47D transcripts could be detected in RNA from human tumor cell lines such as melanoma (S361), lung cancer (A549), colorectal adenocarcinoma (SW480), cervix carcinoma (HeLa), Burkitt’s lymphoma (Raji), lymphoblastic leukemia (Molt-4), promyelocytic leukemia (HL-60), and chronic myeloid leukemia (K562) and in RNA from peripheral blood mononuclear cells of patients with chronic myeloid leukemia, acute lymphatic leukemia, or acute myeloid leukemia (data not shown). These results suggest that HERV-K-T47D transcription at levels detectable by Northern blot analysis is tissue specific and steroid hormone dependent.

**Transcriptional activity of the 5’ LTR of HERV-K-T47D.** To examine the promoter activity of the putative 5’ LTR of HERV-K-T47D in T47D cells, plasmids containing the luciferase reporter gene downstream of an LTR-containing DNA sequence by using forward primer GCGGGATCCGAGGCAAGGAGCTGCAAGGATTCTAAG, nt 1358 to 1381), 

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To confirm our Northern blotting results and further analyze the observed truncated HERV-K-T47D transcript, PCR experiments were performed on T47D cDNA, using primers derived from HERV-K-T47D gag (reverse primer, CGCGGATCCCTACAGTTGGAAACCAAGGGC (nt 1221 to 1243). Employing the newly introduced BamHI restriction sites, BB1.2 was cloned in both the sense (pB-L-BB1.2s) and the antisense (pB-L-BB1.2as) directions into the multiple cloning site of the luciferase expression vector pB-L (Fig. 5A) (kindly provided by H.-J. Butterfass, German Cancer Research Center, Heidelberg, Germany), or the human ubiquitin gene (Clontech) was used to monitor mRNA integrity. Hybridization with the HERV-K-T47D pol fragment under high-stringency conditions revealed a high level of transcription exclusively in full-term placental tissue (Fig. 4B). The signal corresponds in size to the band obtained from T47D RNA after steroid treatment (Fig. 4A). With the exception of the breast carcinoma cell line T47D, a correlation between HERV-K-T47D expression and the occurrence or progression of malignancy was not found. No HERV-K-T47D transcripts could be detected in RNA from human tumor cell lines such as melanoma (S361), lung cancer (A549), colorectal adenocarcinoma (SW480), cervix carcinoma (HeLa), Burkitt’s lymphoma (Raji), lymphoblastic leukemia (Molt-4), promyelocytic leukemia (HL-60), and chronic myeloid leukemia (K562) and in RNA from peripheral blood mononuclear cells of patients with chronic myeloid leukemia, acute lymphatic leukemia, or acute myeloid leukemia (data not shown). These results suggest that HERV-K-T47D transcription at levels detectable by Northern blot analysis is tissue specific and steroid hormone dependent.

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used for internal standardization). The total amount of DNA per dish was adjusted to 6.5 μg with pBluescript SK(+) vector. Incubation was carried out for 16 to 18 h at 37°C and 5% CO₂. T47D cells were then further incubated in fresh RPMI 1640 medium for 48 h. T47D cells were treated with estradiol and then progesterone (in dimethyl sulfoxide solvent), each for 24 h, as described by Keydar et al. (11), while the control dishes received medium with dimethyl sulfoxide alone. At 48 h postincubation (37°C, 5% CO₂), cells were harvested and lysates were prepared according to the recommendations of the Enhanced Luciferase Assay Kit (Berthold Detection Systems, Pforzheim, Germany). Relative HERV promoter activity was calculated as the ratio between the levels of luciferase expression of the constructs and the pBl vector. Transient expression of the constructs in T47D cells revealed that pBL-BB1.2s displayed about the same relative transcriptional activity as the active promoter of the HERV-H LTR (pBL-HERV-H). Upon steroid induction of T47D cells, an about twofold enhancement of luciferase activity was observed (data not shown). This suggests that the 5’ LTR of HERV-K-T47D contains regulatory elements that are steroid dependent and can mediate efficient transcriptional activity of HERV-K-T47D or other sequences in T47D cells.

In conclusion, our results show that HERV-K-T47D is actively transcribed in T47D cells in a steroid-dependent manner, and this active transcription is easily accounted for by the promoter activity and the presence of a number of putative transcription factor binding sequences found in the 5’ LTR. Such activity may also apply to a number of related solitary LTRs which were also detected, perhaps resulting in transcriptional activation of disease-associated antigens. However, the HERV-K-T47D-specific transcript, containing only gag and pol sequences, does not comprise a full-length proviral sequence but is presumably irregularly spliced or terminated. Since HERV-K-T47D does not have the coding capacity for full-length structural proteins, the origin of the retroviral proteins responsible for particle formation and the RT activity found associated with T47D particles (11, 32, 38) is still unclear. Particularly, the gag gene, which is essential for virus packaging and particle formation, is inactivated by stop codons and frameshifts in HERV-K-T47D. As rescue experiments with defective retroviruses lacking the gag, pol, and env open reading frames suggest (41), these activities must be provided in trans by other coding-competent HERV elements. Examples of such coding-competent HERVs are members of the HERV-K(HML-2) subgroup, transcripts of which have been detected in some human teratocarcinoma cell lines (20, 42). Since we were not able to isolate intact protein-coding HERVs from particle preparations, the packaging signals of these genomes may be defective in order to prevent the generation of replication-competent and possibly infective retroviral particles. Particularly, in light of the use of retroviral vectors in gene therapy or the prospect of xenotransplantation (40, 43), identification of such HERV sequences and understanding the mechanisms and risks of generating new, infectious retroviral particles will be of major importance.

Nucleotide sequence accession number. The complete nucleotide sequence of HERV-K-T47D has been deposited in GenBank under accession no. AF020092.

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FIG. 5. Analysis of HERV-K-T47D putative LTR promoter activity in T47D cells. (A) pBL-HERV reporter constructs used for luciferase expression assays. The putative LTR 1.2-kb PCR fragment (Fig. 1, fragment BB1.2) was cloned in the sense (pBL-BB1.2s) and the antisense (pBL-BB1.2as) orientations into the luciferase expression vector pBL. As controls, plasmid pBL-SH1.5gag with the insert (SH1.5gag, Fig. 1) and pBL-HERV-H containing the HERV-H LTR promoter of H6 (7) were similarly constructed. MCS, multiple cloning site; SV-40, simian virus 40. (B) Transient expression in T47D cells of HERV-pBL luciferase reporter constructs. T47D cells were transiently transfected according to standard procedures. The luciferase expression driven by the retroviral promoter was measured by a standardized luciferase assay and is shown as bar graphs representing relative promoter activity. All results shown are derived from triplicate experiments.
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