A Non-Long Terminal Repeat Retrotransposon Family Is Restricted to the Germ Line Micronucleus of the Ciliated Protozoan *Tetrahymena thermophila*

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The ciliated protozoan *Tetrahymena thermophila* undergoes extensive programmed DNA rearrangements during the development of a somatic macronucleus from the germ line micronucleus in its sexual cycle. To investigate the relationship between programmed DNA rearrangements and transposable elements, we identified several members of a family of non-long terminal repeat (LTR) retrotransposons (retroposons) in *T. thermophila*, the first characterized in the ciliated protozoa. This multiple-copy retrotransposon family is restricted to the micronucleus of *T. thermophila*. The REP (*Tetrahymena* non-LTR retroposon) elements encode an ORF2 typical of non-LTR elements that contains apurinic/apyrimidinic endonuclease (APE) and reverse transcriptase (RT) domains. Phylogenetic analysis of the RT and APE domains indicates that the element forms a deep-branching clade within the non-LTR retrotransposon family. Northern analysis with a probe to the conserved RT domain indicates that transcripts from the element are small and heterogeneous in length during early macronuclear development. The presence of a repeated transposable element in the genome is consistent with the model that programmed DNA deletion in *T. thermophila* evolved as a method of eliminating deleterious transposons from the somatic macronucleus.

Developmentally programmed DNA rearrangements occur in a wide variety of organisms (reviewed in reference 5). Functions such as altering gene dosage or directly regulating gene expression have been assigned to many but not all examples of programmed DNA rearrangements. A clinically important example of a programmed DNA rearrangement is V(D)J recombination (2). In addition, a variety of mammalian parasites use programmed DNA rearrangements to vary their surface antigens to avoid host immune response (4). The function of other programmed DNA rearrangements is not as clear. The extensive genome rearrangements that occur during nuclear development in the ciliated protozoa provide an example of programmed DNA rearrangements with poorly understood function.

Like all ciliated protozoa, the oligohymenophoran *Tetrahymena thermophila* displays nuclear dimorphism with a mostly transcriptionally silent diploid germ line nucleus (micronucleus) and a polyploid and transcriptionally active somatic nucleus (macronucleus) within the same cell. The macronucleus develops from a mitotic product of the micronucleus during conjugation. When two cells of different mating types conjugate, the micronucleus in each divides meiotically and mitotically to generate a haploid gametic nucleus that is reciprocally exchanged and fuses with that of its partner to form a zygotic nucleus. This zygotic nucleus divides and from one of the products develops a new macronucleus, while the old macronucleus is concurrently degraded. In *T. thermophila*, macronuclear development involves extensive programmed DNA rearrangements, including chromosome fragmentation, DNA amplification, and site-specific interstitial DNA deletion (12). Interstitial DNA deletion is responsible for the elimination of approximately 10 to 15% of the germ line genome representing more than 5,000 single and multicopy elements. The size of these internal eliminated sequences in *T. thermophila* ranges from 0.6 kb to over 22 kb. Internal eliminated sequences have not yet been found in the coding sequence in *T. thermophila*, but two are located within introns (11, 24).

One model describing the possible function of micronucleus-limited DNA and possible reasons for its absence from the macronucleus proposes that it has a specific function in the germ line micronucleus, serving as centromeres or matrix attachment regions, for example (12, 26). Another model (32) suggests that micronucleus-limited DNA elements represent the remains of a transposon invasion and are eliminated from the streamlined, somatic nucleus to prevent the elimination of essential genes. This model suggests that internal eliminated sequences retain only the *cis* elements necessary for recognition by a *trans*-acting excision machinery provided by the host genome. These models are not necessarily mutually exclusive: repetitive DNA families including transposable elements are often found in centromeric or telomeric regions in higher organisms (35, 43).

The finding that small RNAs, the *TWI1* gene product, and histone methylation have roles in DNA elimination in *T. thermophila* (47, 52) provides a possible molecular link between transposable elements and programmed DNA rearrangements. *Twi1p* belongs to the PPD family, a group of proteins containing conserved Piwi and PAZ domains, some members...
of which are involved in RNA interference as well as the silencing of transposable elements (51). These observations led to the development of the scan RNA model of DNA elimination (47) that suggests that small RNAs are used by the cell to “template” regions of DNA for elimination.

Transposable elements may be divided into two general classes, DNA transposons and RNA-based retrotransposons. Retrotransposons transpose via an RNA intermediate mediated by a reverse transcriptase (RT) (reviewed in reference 16). Retrotransposons are classified into two main groups. The first group of elements are flanked by long terminal repeats (LTRs) and transpose through a double-stranded DNA intermediate synthesized from the RNA template and then integrate with a mechanism similar to that used by DNA transposons (reviewed in reference 16). LTR retrotransposons contain gag and pol genes and are structurally similar to retroviruses, such as human immunodeficiency virus. Non-LTR elements (retroposons) lack terminal repeats and transpose by a mechanism termed target-primed reverse transcription, reverse transcribing a cDNA copy of their RNA directly into the nicked chromosomal target site (36). Non-LTR retrotransposons (also called long interspersed nuclear elements) are widespread in eukaryotes but absent from the model yeast Saccharomyces cerevisiae.

There are several examples of non-LTR retrotransposons with potential specific chromosomal function. The TART and HeT-A non-LTR retrotransposons transpose to the ends of the chromosomes and compensate for the absence of canonical telomeres in Drosophila melanogaster (35). In Giardia lamblia two families of non-LTR retrotransposons are found exclusively at the ends of chromosomes in head-to-tail arrays, where they may form a buffer between telomeric sequence and single-copy genes (1). In humans, the Lyon repeat hypothesis postulates that long interspersed nuclear elements play a role in the developmentally regulated silencing of the female X chromosome (37, 38).

DNA-based transposons have been characterized in the micronuclear genome of several spirotrichous ciliates (14, 28–30). In T. thermophila, several transposon-like families have been identified. The Tel-1 family is restricted to the micronucleus and structurally resembles transposons (10). The Tel-1 elements are found in the micronucleus, adjacent to internal blocks of tandemly repeated 40-bp telomeric repeat and do not contain open reading frames (ORFs). The Tel family is also micronuclear limited (54), with the best-characterized family member, Tel1, containing an 825-bp terminal inverted repeat separated by a long (22-kb) internal region (56). The internal conserved region contains numerous ORFs, some transposon-like, including a retrovirus-like integrase (22), and several that appear to be viral in origin (56). Sequences flanking most Tel and Tel-1 elements are micronuclear limited, suggesting that they are eliminated as parts of larger tracts of micronuclear-limited sequence (56). Although there has been some speculation that the Tel family could transpose through an RNA intermediate (22), no canonical retrotransposons have been found to date in the ciliated protozoa.

Repeated sequences that are retrotransposon-like in origin comprise large parts of the genomes of higher organisms (13) but have not yet been identified in the ciliated protozoa. To investigate possible links between retrotransposons and genome rearrangements in the ciliated protozoa, we identified a family of non-LTR elements in T. thermophila. The REP (Tetrahymena non-LTR retrotransposon) elements appear to be phylogenetically isolated, consistent with the idea that non-LTR retrotransposons are transmitted vertically (16). Consistent with its micronucleus-limited position and the scan RNA model for DNA elimination in T. thermophila (47), REP element transcripts during early conjugation, including meiosis, are heterogeneous in size. The presence of a repeated transposable element within the micronuclear genome is consistent with the model that internal eliminated sequence excision in T. thermophila evolved as a method of ridding the somatic macronucleus of transposable elements.

**Materials and Methods**

**Cell strains.** T. thermophila strains CU428 [Mpr+/Mpr− (VII, mp-s)] and B2086 [Mpr+/Mpr− (II, mp-s)] of inbreeding line B were provided by J. Gaertig, University of Georgia, Athens. Cells were cultured as previously described (19).

**Isolation of micro- and macronuclei.** Micro- and macronuclei were isolated from strain CU428, grown to mid-log phase in 1 × SPP (19) by the method of Gorovsky et al. (23) as modified by Howard and Blackburn (27), without the filtration step.

**Library screening.** The pMBR micronuclear genomic library (provided by Kathleen Karrer, Marquette University, Milwaukee, Wis.) consists of partially MboI-digested micronuclear DNA from T. thermophila cloned into the SacI site of pUC19 (49). An aliquot (100 ng) of library DNA was transformed into Escherichia coli DH5αF', made competent with rubidium chloride (50). The library was screened with standard procedures (50). Probes were gel-purified and labeled by random priming (50) with [α-32P]dATP (Amersham).

**Southern blotting and DNA manipulations.** Southern molecular biology techniques were performed as described by Sambrook et al. (50) or by following supplier’s instructions. DNA-modifying enzymes were obtained from New England Biolabs. Southern blots were hybridized in 1% sodium dodecyl sulfate (SDS)–5× SSC–0.5% dry milk powder and washed three times at 20 min each in 1× SSC–0.1% SDS, once in 0.1× SSC–0.1% SDS at 30°C, and imaged with either X-ray film or a Canberra Packard Instant Imager.

**RNA isolation and Northern analysis.** Total cell RNA from T. thermophila was isolated with the Trizol reagent (BRL); 6 × 10⁶ cells were concentrated into 100 μl, to which 1 ml of Trizol was added. Whole-cell RNA was purified according to instructions provided by the supplier. The RNA pellet was air-dried before being dissolved in RNase-free double-distilled H₂O at 4 μg/μl. Northern analysis was performed by separating 20 μg of total RNA on a 1% agarose–0.66 M formaldehyde gel. RNA was transferred to a nylon filter after three 20-min washes in 10× SSC (50). Prehybridization and hybridization were performed at 42°C in 50% formamide–5× Denhardt’s solution–1% SDS–5× SSC with sheared, denatured salmon sperm DNA at 100 μg/ml. Filters were washed three times for 15 min in 1× SSC–0.1% SDS at room temperature, and 1 time in 0.25× SSC–0.1% SDS at 42°C for 15 min after overnight incubation with the indicated probes. Filters were stripped of hybridized probe by boiling in 1× TE–1% SDS for 5 min, cooling to room temperature over 10 min, and then washing in 5× SSC for 5 min. The stripped filters were immediately covered with Saran Wrap before being imaged for verification and then stored at 4°C.

**Table 1. Sequences of oligonucleotides used for PCR amplification in this study**

<table>
<thead>
<tr>
<th>Oligonucleotide</th>
<th>Sequence</th>
</tr>
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<tbody>
<tr>
<td>TTRTR</td>
<td>5'-GGTTGCTGAAGGCTACTATAG</td>
</tr>
<tr>
<td>RTFII</td>
<td>5’-GAACAAAACTTAAACGGATATT</td>
</tr>
<tr>
<td>TIPK</td>
<td>5’-GARATHCCHGGGYARTAY</td>
</tr>
<tr>
<td>TPKR</td>
<td>5’-GYDGTNARHGYRAADDG</td>
</tr>
<tr>
<td>RTR</td>
<td>5’-GATTGCTGAAGGCTACG</td>
</tr>
<tr>
<td>REV1</td>
<td>5’-GAATTCTGAGTGAGGATAGTGA</td>
</tr>
<tr>
<td>FOR3</td>
<td>5’-TGTACGTTCTTGTGAAG</td>
</tr>
</tbody>
</table>

*R = A + G; H = A + T + C; N = A + T + C + G; Y = T + C; D = A + T + G.*
Oligonucleotides. See Table 1 for the oligonucleotides used during this study.

PCR conditions. PCR was performed as described previously (19). Long-range PCR was performed with the Expand Long Template PCR system (Boehringer Mannheim) with conditions as specified by the supplier. To obtain 5' H11032 sequence from REP6, we first used inverse PCR with primers targeted within the macronucleus-destined HEH2 gene (Fig. 1) to amplify and then sequence the macronuclear locus of the region. We then amplified the 5' H11032 end of REP6 with a primer in the 5' H11032 macronuclear-destined sequence (FOR3, Table 1) and one in REP6 (REV1, Table 1) with whole-cell DNA as a template.

DNA sequencing. Sequencing was performed with automated cycle sequencing with dye-labeled dideoxy terminators and a PE/ABI 373a or 377 sequencer at the Core Molecular Biology Facility, York University, Toronto, Ontario, Canada.

Sequence assembly and analysis. The pMBR clones were separately subjected to transposon-based random insertion with the GPS-1 genome priming system (New England Biolabs, Mississauga, Ontario). Restriction enzyme mapping was used to identify a series of subclones that when individually sequenced with universal primers provided for the GPS-1 kit for pMBR-RP1, pMBR-RP2, pMBR-RP4, and pMBR-RP6 were assembled by pairwise sequence alignments with BlastN (http://www.ncbi.nlm.nih.gov/blast/bl2seq/bl2.html). Gaps were filled by DNA sequencing with specifically designed oligonucleotides. When the entire sequence across each plasmid insert was obtained, it was scanned for open reading frames (ORFs) with homology to known proteins by searching the nonredundant protein databases with the BlastX algorithm. DNA sequences identified as containing a possible match to a known protein were translated with the ciliate genetic code and used to query the nonredundant protein databases with the BlastP algorithm. Reverse position-specific Blast was used to search these regions for potential conserved domains with the conserved domain database of NCBI (http://www.ncbi.nlm.nih.gov/Structure/cdd/wrpsb.cgi). The Simple Modular Architecture Research Tool (SMART: http://smart.embl-heidelberg.de/) was also utilized for this purpose.

Multiple sequence alignments of the reverse transcriptase (RT) and apurinic/apyrimidinic endonuclease (APE) domains were performed by sequentially adding the sequence of the APE and RT domains of ORF2 from the T. thermophila REP elements from the pMBR-RP clones to the APE and RT profile alignments of Malik et al. (40) with the profile alignment option of ClustalW (http://bioweb.pasteur.fr/seqanal/interfaces/clustalw.html#profile), where profile 1 was either accession number DS36736 [APE domain (40)] or ds36752 [RT domain (40)] (ftp://ftp.ebi.ac.uk/pub/databases/embl/align/) and profile 2 was the T. thermophila REP element sequences. Alignments were adjusted by hand and smaller versions were shaded by importing the ALN file into the Boxshade server (http://www.ch.embnet.org/software/BOX_form.html) and then editing the postscript file in Adobe Illustrator 9.0.

Phylogenetic analysis of the RT and APE domains. The resulting profile alignments were used to construct a neighbor-joining tree (N = 1,000 replicates) with ClustalW (http://bioweb.pasteur.fr/seqanal/interfaces/clustalw.html#trees) which was visualized with the Phylodendron Phylogenetic tree printer (http://www.es.embnet.org/Software/BOX_form.html) and then editing the postscript file in Adobe Illustrator 9.0.
Nucleotide sequence accession numbers. GenBank accession numbers for the DNA sequences of pMBR-RP1 to pMBR-RP3 and pMBR-RP6 are AY371728 to AY371731, respectively.

RESULTS

*T. thermophila* contains a non-LTR retrotransposon family.

We used degenerate primers (TTPKF and TTPKR, Table 1) to amplify from *T. thermophila* whole-cell DNA a sequence of ~700 bp that, when compared against the nonredundant NCBI database, yielded similarity to the RT domain of non-LTR retrotransposons. To amplify DNA sequence containing the RT domain, we used inverse PCR (48) of HindIII-digested *T. thermophila* whole-cell DNA with primers RTF and RTR (Table 1) to obtain 2.2 kb of sequence, a portion of which was used as a DNA probe to screen a *T. thermophila* micronuclear DNA library. Hybridizing clones were abundant, suggesting that the sequence is repetitive in the *T. thermophila* micronuclear genome. Four independent clones were sequenced in full. Pairwise and multiple sequence alignments of the four clones revealed regions of extensive DNA sequence similarity. Further sequence analysis revealed that the similarity arises due to the presence in each plasmid of either a full or partial non-LTR retrotransposon.

Partial restriction maps showing the micronuclear genomic structure of the four pMBR-clones are presented in Fig. 1. Plasmids pMBR-RP1 and -RP3 each contain partial sequence of one REP element (named REP1 and REP3, respectively; see boxed regions, Fig. 1). Plasmid pMBR-RP2 contains one full REP element (REP2-2) as well as partial sequence of a second REP element (REP2-1; see boxed region, Fig. 1). In addition, plasmid pMBR-RP6 contains partial sequence of an additional REP element (REP6; Fig. 1). We amplified additional DNA sequence from *T. thermophila* micronuclear DNA to complete the entire REP6 sequence (Fig. 1).

5′UTR of the REP elements. The REP2-2 element is the only REP element identified within the four plasmids potentially to contain an entire REP element with an intact 5′ untranslated region (5′UTR). We verified the 5′ limit of REP element sequence with the additional amplified REP6 sequence (see Fig. 1 and Materials and Methods). We aligned REP2-2 and the complete REP6 sequence and identified a 243-bp region immediately upstream of ORF1 (Fig. 1) that is highly conserved in both REP6 and REP2-2 and is therefore predicted to comprise the entire 5′UTR region of the REP element. The upstream boundary of REP2-2 is therefore the predicted 3′UTR and/or flanking genomic sequence of REP2-1 (Fig. 1).

**ORF1.** A potential ORF1 in REP elements was detected by translating the nucleotide sequence upstream of the previously identified ORF2 of REP1 and REP2-2 (Fig. 1) and aligning the predicted protein sequences. The 412-amino-acid protein conceptually encoded by ORF1 of REP2-2 does not contain any significant similarity to other proteins. REP1 contains the DNA encoding the C-terminal 368 amino acids of another ORF1. The overlapping regions of the two ORF1s are 95% identical at the amino acid level (see Fig. A in supplemental material, http://www.yorku.ca/ronp/). Analysis of the predicted amino acid sequence with SMART indicates the possible presence of a zinc finger related to the CCCH class (Fig. B in supplemental material, http://www.yorku.ca/ronp/) as well as a coiled-coil region. The CCCH class of the zinc finger superfamily is found in many RNA-binding proteins and, to our knowledge, has not been found in any non-LTR retrotransposons described to date.

**ORF2.** The main conserved feature that the REP elements share with other non-LTR retrotransposons is the 1,147-amino-acid ORF2. Conserved domain analysis of the ORF2 amino acid sequence identified two protein domains characteristic of non-LTR retrotransposons: a reverse transcriptase (RT) domain, and an apurinic/apyrimidinic endonuclease (APE) domain. Many non-LTR retrotransposons encode an APE domain which is related to the AP endonuclease that function in DNA repair (40, 44). The APE domain of at least one non-LTR retrotransposon has been shown to nick DNA with sequence specificity related to its target site (18), although not all non-LTR retrotransposons are associated with specific genomic target sites (40). A multiple sequence alignment of the REP APE domain with that of several other non-LTR retrotransposons is shown in Fig. 2. The REP APE is likely to encode a functional endonuclease, as there is excellent conservation of key residues that have been identified in the APE active site (45).

Structurally, RT enzymes adopt a hand structure that has a palm containing the active site and fingers and a thumb that loosely hold the RNA template while the RT moves laterally. Xiong and Eickbush (58) found that there are seven blocks of amino acid identity common to the RT of all retroelements. The REP family contains the seven classic domains 1 to 7, as is expected for a non-LTR retrotransposon (Fig. 3). Burke et al. (6) extended non-LTR RT homology to 11 blocks of identity (including domains 0, 2a, 8, and 9) by comparing the three-dimensional structure of the RT domain of the retrovirus human immunodeficiency virus (33) to the RT domain of the R2 non-LTR retrotransposon. The highly conserved PGPD motif in domain 0 (Fig. 3) is absent in the REP family. Burke et al. (6) suggest that this domain is believed, in combination with domain 2a, which is present in the REP elements (Fig. 3), to extend the active site of the non-LTR RT by giving additional or longer fingers compared with that of human immunodeficiency virus RT.

Domain 8 appears to be relatively conserved between the REP elements and non-LTR retrotransposons (Fig. 3). Domains 9 and 9* have been used to divide non-LTR transposons into two subgroups (Fig. 3) based upon the presence or absence of a glycine-rich tetramer (40). The multiple sequence alignment (Fig. 3) places the REP elements into the subgroup without the glycine-rich tetramer (domain 9, Fig. 3). It is believed that the function of domain 8 and 9 is to extend the thumb. Domain 7 was identified as the thumb region of human immunodeficiency virus RT (33). It has recently been suggested that the presence of domains 0 and 2a (more/longer fingers) and 8 and 9 (extended thumb) allows the RT to more completely wrap around its RNA template. The biological relevance of this would be higher processivity of the non-LTR RT compared to those of retroviruses and LTR retrotransposons (3). The biological relevance of the absent domain 0 in the REP element RT is not known. The ORF2 of the *T. thermophila* REP element does not appear to encode a C-terminal DNA-binding domain or an RNase H domain similar to human long interspersed nuclear elements.

**3′UTR of REP elements.** Pairwise or multiple sequence alignments of the sequence immediately downstream of the
Stop codon of each of the five identified full or partial REP elements do not show sequence identity except that four out of the five immediately follow the stop codon with the sequence 5'-AAT/GAA-3'. Four of the five REP elements contain two to eight copies of different tandemly repeated DNA sequence (Fig. C in supplemental material, http://www.biol.yorku.ca/ronp/). The REP element of pMBR-RP3 does not appear to have this sequence organization (Fig. 1). The only conserved DNA sequence common to all five REP elements is a highly conserved 54-bp sequence that is found a distance of 404 bp to 932 bp from the respective REP element stop codon (Fig. 4).

From the analysis of REP2-2 and REP6, we estimate that the size of an intact REP element is approximately 6.1 kb. Downstream of the three imperfect repeats that flank the REP2-2 element is a region that contains a sequence that appears to be the C terminus of an ORF encoded by a DNA transposon of the Tc-1 family. Only partial sequence of the putative Tc-like ORF is available as the 3' boundary of pMBR-RP2 interrupts the putative coding region of this DNA transposon (Fig. 1). The putative T. thermophila Tc-like element is organized in opposite orientation to the two REP elements (Fig. 1).

The 5' end of pMBR-RP3 contains the C-terminal 508 amino acids of ORF2 of the pMBR-RP3 REP element. The 350-bp sequence that flanks the REP1 and both REP2 elements does not similarly flank REP3. No significant matches were found when the pMBR-RP3 sequence downstream of the conserved 54-bp sequence was used in a BlastX search against the public protein database.

The 5' end of pMBR-RP6 contains the C-terminal 820 amino acids of the REP6 ORF2. The pMBR-RP6 sequence downstream of the conserved 54-bp sequence (Fig. 1) contains high similarity to human KIAA1279 (31), a gene of unknown function expressed in the human brain. This gene (HEH2) is expressed during vegetative growth of T. thermophila and its protein product has been localized to basal bodies (21). We used Southern blotting and PCR to determine that the HEH2 gene is macronuclearly retained (data not shown). The REP6 locus has been mapped to the right arm of micronuclear chromosome 2 (Hamilton and Orias, personal communication).

REP elements are restricted to the germ line micronucleus. The possible relationship between transposable elements and...
FIG. 3. Multiple sequence alignment of the RT domain of REP1, REP2-2, REP3, and REP6 with that of several non-LTR retrotransposons (see Materials and Methods for details). RT subdomains are indicated by a corresponding number above the conserved block of amino acid sequence (see text for details). With respect to domain 9, non-LTR retrotransposons are divided into two subgroups based upon the presence or absence of a glycine-rich tetramer (domain 9 versus 9*: see boxed region). Shading of amino acid residues is as in Fig. 2.
programmed DNA elimination in the ciliated protozoa led us to predict that REP elements would not be found in the somatic macronucleus. Southern analysis of DNA from macronuclei and micronuclei probed with the conserved RT domain showed strong hybridization to micronuclear but not to macronuclear DNA (Fig. 6).

Nucleotide substitutions in ORF2 are nonrandom with respect to codon position. Nucleotide substitutions in several of the open reading frames of the transposon-like Tlr elements are nonrandom with respect to their codon position, with the majority occurring in the third position (22, 56), implying that the ORFs are under selective pressure to preserve their function. We analyzed two regions of ORF2 of the identified REP elements for codon position of nucleotide changes: the APE domain of REP1 and REP2-2, and the RT domain of REP1, REP2-2, REP3, and REP6. If the REP elements were not under selective pressure to maintain a functional protein, nucleotide changes should occur at random with an equal frequency at each position. According to $\chi^2$ analysis, nucleotide changes are nonrandom with respect to codon position ($P < 0.001$), with the majority occurring in the third position (Table 2). In agreement with this result, considering only the RT domain, 93% of the identified 215-nucleotide polymorphisms (Table 2) lead to either identical or similar amino acids, as defined by the Structure-Genetic matrix scoring system (17). Although the sample size of the APE domain is smaller, the result is striking, with 100% of nucleotide polymorphisms leading to identical or similar amino acids (Table 2).

The noncanonical genetic code of *T. thermophila* (25) dictates that the “universal” stop codons TAA and TAG encode glutamine (Q), ensuring that $T \rightarrow C$ transitions in the first nucleotide of TAA or TAG codons are silent. For the RT domain, 11 out of 16, or 69%, nucleotide changes resulting in the same amino acid were one of these two transitions. A similar result, 11 out of 18, or 60%, was obtained when the entire ORF2 of REP1 and REP2-2 was considered.

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**FIG. 4.** Multiple sequence alignment of the conserved 54-bp sequence found 3' of ORF2 of the REP elements. An arrow represents the point within the conserved sequence where there is a small amount of overlap with the repetitive sequence described in Fig. 5. The numbering begins at the stop codon of ORF2 of the respective REP element. The black shading indicates nucleotides that are identical in all five sequences.

**FIG. 5.** Multiple sequence alignment of the ~350-bp imperfectly repeated sequence found immediately downstream of the conserved 54-mer (Fig. 4; included here as a part of REP1, REP2-1, and REP2-2-1). The sequence similarity between the repeats of REP2-1, REP2-2-1, and REP2-2-2 extends further than that of REP1 and REP2-2-3. Nucleotide residues that are identical in all five sequences are shaded black.
Phylogeny of the *T. thermophila* REP elements. The above analysis indicates that the REP elements have evolved within the context of the *T. thermophila* genome. Since non-LTR retrotransposons are primarily transmitted vertically (40), and to date no non-LTR retrotransposons have been identified in the ciliated protozoa, we would predict that REP elements should be grouped apart in molecular phylogenies of the non-LTR retrotransposon family. We aligned separately the RT domains (REP1, REP2-2, and REP3) and the APE domains (REP1 and REP2-2) with the respective domains from non-LTR retrotransposons (40) deposited in the EMBL database (Fig. 2 and 3 and Materials and Methods). Possible phylogenetic relationships inferred from neighbor-joining trees constructed from the alignments of the RT and the APE domains are presented in Fig. 7 (RT) and Fig. 8 (APE).

Malik et al. (40) proposed the term clade to represent those non-LTR retrotransposon elements that are grouped together with some other small differences in this phylogeny, such as the RT clade placed within the L1 clade (Fig. 8) and LOA within the R1 clade [(Fig. 8; also seen in Malik et al. (40)], although the bootstrap values at the nodes of these long-branching divergences are not high. This may be due to the fact that the APE phylogeny has less resolution than the RT phylogeny (40).

We conclude that the *T. thermophila* REP elements represent a unique clade of non-LTR retrotransposons. The grouping of the APE domain of the REP elements with that of the DRE element may signify that the APE domain of the REP elements is evolving under tight functional constraints. Consistent with this hypothesis, we did not detect nucleotide changes that resulted in dissimilar amino acids in the analysis of the APE domains of REP1 and REP2-2 (Table 2).

REP elements are transcribed during nuclear development in *T. thermophila*. To analyze transcription of the REP elements during growth or nuclear development, we isolated whole cell RNA from growing, starved or conjugating *T. thermophila* and analyzed it by Northern blotting, probing with the RT of the REP element (Fig. 9). During early conjugation, we observed heterogeneously sized transcripts that ranged in size from ~0.5 kb to 1.0 kb. They were visible as early as 3 h after the initiation of conjugation, a time corresponding to micronuclear meiosis (46).

**DISCUSSION**

The REP elements of *T. thermophila* represent a new clade of non-LTR retrotransposons. We have characterized the first non-LTR retrotransposon from the ciliated protozoa. Similar to all transposable elements identified to date in the ciliates, the REP family appears to be restricted to the micronucleus. Phylogenetic analysis suggests that the REP family is relatively divergent compared to identified non-LTR retrotransposon clades (40). This is consistent with the results of our analysis of the nature of nucleotide changes in the ORF2 that suggest that the REP elements have evolved in the context of the genome of *T. thermophila* (Table 2). These data are also consistent with the fact that maintenance of non-LTR retrotransposons in the genome of a particular organism is almost exclusively by vertical and not horizontal transmission (16).

**Structure of the *T. thermophila* REP element.** The REP element shares many features with the non-LTR retrotransposon family. Many non-LTR retrotransposons encode an N-terminal ORF1 protein, most of which contain CCHC zinc finger motifs that are possibly involved in binding the element RNA after translation to import it back into the nucleus for transposition. The REP elements encode a putative ORF1 that contains an unusual CCHC zinc finger motif. The fact that some CCHC zinc finger-containing proteins have been shown...
FIG. 7. Phylogenetic relationships of the *T. thermophila* REP element with other non-LTR retrotransposons based upon the RT domain as defined in Malik et al. (40). The tree is neighbor joining with bootstrap values indicated as number out of 1,000. Only bootstrap values of >50% are indicated. The amino acid divergence scale is indicated. The alignment and amino acid sequences were derived as described in Materials and Methods. The non-LTR elements are divided into clades based upon phylogenetic relationships. Clades are indicated by the black bars and are defined in Malik et al. (40).
to specifically bind 3'UTR mRNA (34) raises the possibility that ORF1 of the T. thermophila REP element has a similar function.

The REP ORF2 contains both an APE and an RT domain. All non-LTR retrotransposons encode an RT domain and an endonuclease which is either an N-terminal APE like the REP elements or one C-terminal of the RT similar to various prokaryotic restriction endonucleases (59). The APE domain of the REP element is predicted to be functional since key residues in the active site are conserved. Except for domain 0, key RT residues in the RT domain are also conserved. The function of this small domain is unknown at this point, but it is extremely well conserved within the non-LTR group of RTs (40). In combination with domain 2a, which is present in the REP elements (Fig. 4), it is believed to extend the active site of the non-LTR RT by giving additional or longer fingers compared with those of retroviral RTs. The conservation of key residues in both the APE and RT domains

FIG. 8. Phylogenetic relationships of the T. thermophila REP element with other non-LTR retrotransposons based upon the APE domain as defined in Malik et al. (40). The tree is neighbor joining with bootstrap values indicated as number out of 1,000. Only bootstrap values of >50% are indicated. The amino acid divergence scale is indicated. The alignment and amino acid sequences were derived as described in Materials and Methods. The non-LTR elements are divided into clades based upon phylogenetic relationships. Clades are indicated by the black bars and as defined in Malik et al. (40).
suggests that ORF2 remains catalytically active despite its phylogenetic age.

The majority of non-LTR retrotransposons contain poly(A) tails in their 3’UTR, the direct result of reverse transcription and transposition of a polyadenylated transcript. Several contain tandem repeats in their 3’UTR (8, 41). These repeats suggest that an abortive cDNA synthesis step by the RT may exist in these non-LTR retrotransposons and this may be mechanistically related to telomere addition (8). The 3’UTR of REP elements is unusual in that there is no evidence for poly(A) tails and four of the five that we have analyzed contain tandem repeats of very different lengths.

Genomic organization of T. thermophila REP elements. One common motif at the 3’ end of the REP elements is a conserved 54-bp sequence. Due to the absence of sequence conservation in the 3’UTR, it is unclear at this point whether this sequence represents a conserved sequence in the REP element itself or forms part of a REP target site for transposition into genomic DNA. Three of the five REP elements are associated with a ~350-bp sequence that appears to be repeated in the micronuclear genome. The fact that two of the REP elements are not associated with this repeat implies that the repeat may be a preferred but not exclusive target site for REP retrotransposition. There are other examples of retrotransposons associated with repetitive sequence. For example, Genie 1 is an ancient lineage of site-specific non-LTR retrotransposons in G. lamblia that is found within a telomeric repeat (7) and centromeric sequence in Arabidopsis thaliana is composed primarily of a repetitive sequence interspersed with retroelements (43).

The REP elements are restricted to the T. thermophila micronucleus. Our preliminary data from Southern blotting experiments have shown that each of the REP elements is flanked by a micronucleus-limited sequence (T. A. Thing, N. Vythilingum, J. S. Fillingham, and R. E. Pearlman, unpublished data). Thus, it is likely that REP elements are eliminated as parts of larger tracts of micronucleus-limited sequence, similar to the Tlr and Tel-1 elements.

T. thermophila REP elements: implication for genome stability. There are several reports of selection acting to limit missense mutations from accumulating in the open reading frames of transposons in various ciliated protozoa (15, 22, 55, 56). We have determined that ORF2 of the T. thermophila REP elements is subject to the same phenomenon. The L1 element, a human non-LTR retrotransposon, has the property of cis preference in that its two ORFs predominantly mobilize only the RNA that encodes it (53). As a non-LTR retrotranspon, the REP elements may also be subject to cis preference, which would dictate that, to remain mobile, a particular transposon must retain its own functional proteins and not rely on them to be provided in trans.

A micronuclear-specific source of RT may have implications for T. thermophila micronucleus genome stability that extend beyond REP element transposition. It has been speculated that the Tlr element could transpose through an RNA intermediate but it does not encode an RT (56). It is possible that the RT of the REP elements could function in cis preference and catalyze the transposition of various sequences in the micronuclear genome.

The restriction of REP elements to the transcriptionally silent micronucleus is consistent with all transposons described to date in the ciliated protozoa. One model to describe the evolution of programmed DNA deletion in the ciliates suggests that it originated to rid the somatic macronucleus of potentially deleterious transposons. This model suggests that nontransposon internal eliminated sequences represent degenerate transposons that have yielded their hypothetical “excisase” in trans, retaining only the necessary cis-acting signals for programmed elimination (32). The recent discovery of a piwi-related gene with a function in developmentally programmed genome rearrangement in T. thermophila (47) supports this model.

The molecular characterization of the piwi gene family has provided a molecular link between the RNA interference phenomenon and genome surveillance (51). The scan RNA model of programmed DNA rearrangements presented by Mochizuki et al. (47) proposes that germ line-specific sequence is transcribed and processed to small “scan RNAs” that are involved in targeting regions of DNA for programmed deletion in T. thermophila. Further evidence in support of this model has been provided by Yao et al. (60), who found that injection of double-stranded RNA into the cell during early conjugation triggers deletion of the targeted genomic regions during macronuclear development. In addition, Wuitschick and Karrer have shown that the multiple-copy Tlr elements contain multiple, redundant cis-acting sequences within their micronucleus-limited sequences that are able to promote programmed DNA deletion, suggesting that Tlr elements are subject to an RNA interference-like system that detects the presence of multiple-copy sequence within the micronucleus (57). The fact that the REP elements are transcribed during early macronuclear development in a manner similar to other micronuclear-limited sequences (9) is consistent with the hypothesis.
that programmed DNA deletion in *T. thermophila* is a method of eliminating transposable elements from the macronucleus. Preliminary dot-blotting analysis indicates that there are between 40 and 175 REP elements per micronucleus (N. Vythi-


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