TEtrimmer: a novel tool to automate the manual curation of transposable elements

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ABSTRACT

Transposable elements (TEs) are repetitive DNA sequences capable of moving within genomes. Accurate annotation and classification of TEs is crucial but challenging due to their sequence diversity and often fragmented occurrence. We present TEtrimmer, a novel tool to automate manual TE curation. TEtrimmer integrates multiple sequence alignment (MSA) clustering, MSA sequence extension, MSA cleaning, TE boundary definition, and TE classification, and provides report plots and a graphical user interface (GUI) application to inspect and improve results. Benchmarked on the genomes of six organisms from various kingdoms of life, TEtrimmer consistently improved the identification of intact TEs compared to established tools.

Keywords: Transposable element; Manual curation; Multiple sequence alignment; Clustering.

Background

Transposable elements (TEs) are selfish repetitive DNA elements that can move within host genomes. TEs were first discovered by Barbara McClintock in maize in 1948 (McClintock, 1950). Since then, TEs have been identified in all studied eukaryotic species where they occupy a large proportion of many genomes, such as around 45% of the human genome (International Human Genome Sequencing Consortium et al., 2001), 53% of the zebrafish genome (Howe et al., 2013), and 85% of the maize genome (Schnable et al., 2009).

TEs were long regarded as "junk" DNA. However, TEs play key roles in genome evolution, development, and immunity (Bourque et al., 2018). For instance, TEs can regulate promoter and enhancer sequences to alter host gene expression (Fueyo et al., 2022). In addition, TEs are a source of many regulatory genes, including long non-coding RNAs (Kapusta et al., 2013) and small RNAs (Kunz et al., 2024). Moreover, TEs can be drivers of evolutionary innovations. A recent example of TE-driven evolution is an *Alu* element insertion into an intron of the human *TBXT* gene, which appears to have contributed to tail-loss evolution in primates (Xia et al., 2024).

TEs are classified according to their transposition mechanism. Class I encompasses the retrotransposons, which copy themselves for genome insertion via an RNA intermediate. Class I retrotransposons mainly include three subclasses, LTR (long terminal repeat) retrotransposons, LINEs (long interspersed nuclear elements), and SINEs (short interspersed nuclear elements) (Bourque et al., 2018; Wicker et al., 2007). An intact LTR retrotransposon always contains flanking terminal repeat sequences, while LINEs and SINEs are defined by their terminal genomic poly-A repeats. Class II, on the other hand, comprises DNA transposons, which do not usually generate copies but rather excise themselves for insertion into another locus in the host genome. The most prominent DNA transposons include TIR (terminal inverted repeat) and Helitron elements (Wicker et al., 2007). However, precise TE annotation remains challenging, mainly due to TE type divergence and the existence of decayed, fragmented, nested, and low-copy TEs, especially in genomes with a high TE content (Hoen et al., 2015; Storer et al., 2022).

TE characterization involves two main steps, which are TE discovery and TE annotation (Storer et al., 2022). Due to the repetitive nature of TEs, TE families can be represented by the TE consensus sequence library, also called TE library, which is constructed based on a multiple sequence alignment (MSA) of the TE copies from the respective reference genome (Goubert et al., 2022). The process of TE library construction is called TE discovery. The TE Hub Consortium (The TE Hub Consortium et al., 2021) has collected 51 tools to date (April 2024) that are associated with TE library construction (Sierra & Durbin, 2024). The term TE annotation refers to the process of genome-wide TE identification to correctly define both intact and fragmented TE copies in the genome. Pre-constructed TE libraries are available for some species from repositories such as RepBase (Weidong. Bao et al., 2015), Dfam (Hubley et al., 2016), and RepetDB (Amselem et al., 2019). To date, RepBase contains the most comprehensive collection of manually curated high-quality TE libraries, but these have been behind a paywall since 2018.

TE annotation is typically based on three main strategies, i.e., repository-based, repeat-based, and structure-based methods. The most-used strategy is the repository-based approach (Loreto et al., 2023), which takes advantage of public pre-constructed TE libraries for whole-genome TE annotation. Since closely related species share a common origin and exhibit genome sequence similarity, the established TE library can be used to identify homologous TEs in related genomes as well (Gao, 2023). This strategy heavily relies on the availability and quality of a suitable TE library for the species of interest. However, manually curated high-quality TE consensus libraries are only available for a limited number of species after years of community efforts (Hubley et al., 2016; Osmanski et al., 2023). In addition,

because the current versions of RepBase are only available via paid subscription, many researchers, particularly from smaller institutes and/or less wealthy countries, do not have access to suitable TE libraries. By contrast, the repeat-based strategy generates a custom TE library from the genome of interest. This strategy mainly includes self-comparison and *k*-mer seeding methods (Storer et al., 2022). The self-comparison method identifies TEs by computationally intensive pairwise alignments between genomic sequences using algorithms such as RECON (Zhirong. Bao & Eddy, 2002) and Grouper (Quesneville et al., 2003). On the other hand, *k*-mer seeding methods like RED (Girgis, 2015), RepeatScout (Price et al., 2005), and P-Clouds (Gu et al., 2008) are generally faster and can efficiently handle large genomic datasets. However, the accurate clustering of TE sequences remains a challenge, and this method relies on high-quality genome assemblies. Poor genome quality can lead to inaccurate TE boundary definition, over-fragmentation of TE copies, and loss of TEs in incomplete assemblies (Kurtz et al., 2008). Finally, the structure-based strategy can annotate TEs by recognizing specific structural hallmarks such as LTRs, TIRs, target site duplications (TSDs), and terminal genomic poly-A repeats. A variety of algorithms, including LTR_FINDER (Xu & Wang, 2007), LTR_STRUC (McCarthy & McDonald, 2003), LTRharvest (Ellinghaus et al., 2008), TIR-Finder (Gambin et al., 2013), and SINE-Finder (Wenke et al., 2011) facilitate structure-based TE annotation. The structure-based methods can efficiently annotate low-copy TEs and precisely define TE boundaries; however, a high false-positive rate is associated with these approaches.

No single strategy or algorithm can comprehensively annotate TEs in a genome. Accordingly, various tools for de novo TE identification have been developed that integrate several annotation strategies, such as RepeatModeler2 (Flynn et al., 2020), EDTA2 (Ou et al., 2019) (https://github.com/oushujun/EDTA/tree/EDTA2), and REPET (Flutre et al., 2011). However, even these tools do not approach the gold standard quality of manually curated TE libraries (Platt et al., 2016).

Manual curation of TEs can be divided into five major steps: BLAST (Basic Local Alignment Search Tool), extend, extract, align, cluster, and trim (Goubert et al., 2022; Storer et al., 2021). In brief, the TE candidate from a de novo discovery pipeline is used for a nucleotide BLAST (BLASTN) query against the reference genome, and the coordinates of search hits are arbitrarily extended at both ends to include TE boundaries. Then, the TE sequences are extracted based on the extended coordinates and subsequently aligned. Finally, the sequences within the MSA are manually clustered based on their sequence relatedness and trimmed by removing regions of low conservation (Goubert et al., 2022; Platt et al., 2016; Storer et al., 2021). The manual curation of TEs can yield a gold standard custom TE library for the genome of interest but is very time-consuming to establish, and requires TE specialists with an in-depth understanding of TE structure and biology (Orozco-Arias et al., 2023). Accordingly, the comparability and reproducibility of TE libraries from manual curation may be limited by the level of expertise of the curators (Baril et al., 2024).

Some software tools, like EarlGrey (Baril et al., 2024) and MCHelper (Orozco-Arias et al., 2023), have been developed to partially automate and assist manual curation of TEs. EarlGrey mainly combines RepeatModeler and TEstrainer (https://github.com/jamesdgalbraith/TEstrainer) for TE library construction. TEstrainer is the core module to automate most manual curation steps but not the clustering of MSAs. TEstrainer attempts to bypass the MSA clustering by only including the top 20 BLASTN hits, which is insufficient for highly divergent TEs. MCHelper includes an MSA clustering method but lacks effective MSA cleaning ability. In essence, both EarlGrey and MCHelper can improve the precision of TE annotation in any given organism but do not yet approach the quality of manually curated TEs. Here, we introduce a novel user-friendly tool called TEtrimmer that automates the manual curation of TEs. TEtrimmer can efficiently cluster MSAs, clean MSAs, and precisely define TE boundaries. TEtrimmer provides comprehensive report plots and a graphical user interface (GUI)

application to support the rapid inspection and improvement of results, which closes the gap to manual curation.

Results

Overview of TEtrimmer

Manual curation is crucial for a high quality of TE annotation. However, curation can involve hundreds to thousands of TE consensus sequences for each genome and is therefore time-consuming and requires expert TE knowledge. TEtrimmer is a software designed to automate the manual curation of TEs. The input of TEtrimmer can be the output TE consensus libraries from de novo TE discovery tools like EDTA2, RepeatModeler2, and REPET and/or TE libraries from closely related species. To allow multi-thread analysis, TEtrimmer separates each set of input sequences into a single file (Fig. 1). For each separated sequence, TEtrimmer automatically performs (BLASTN searches against the respective genome (Ye et al., 2006). It uses the retrieved BLAST hits for MSA generation, MSA clustering (Fig. 2), MSA sequence extension, and MSA cleaning (Fig. 3).

Thereafter, TEtrimmer employs the clustered, extended and cleaned MSAs to generate consensus sequences for the definition of putative TE boundaries. Then, potential terminal repeats are identified, and a prediction of open reading frames (ORFs) and protein domains on the basis of the protein families database (PFAM) (Mistry et al., 2021) are conducted. Subsequently, TE sequences are classified and an output evaluation is performed mainly based on the existence of terminal repeats, and the full length BLASTN hit numbers (Table 1). Consequently, two TE consensus library files are produced by TEtrimmer. The first file contains all output sequences while the second file is generated by removing duplicated sequences via CD-HIT-EST (Fu et al., 2012) (Fig. 1; see Methods for further details).

TEtrimmer provides a series of report plots for each output sequence to enable detailed inspection (Fig. 4). Users can choose to use the TEtrimmer-generated TE consensus library directly for genome-wide TE annotation or take advantage of the supplied GUI application (Fig. 5) to inspect and curate the TE library, which can generate a gold standard manual curation-level TE library. A detailed description of each step can be found in the Methods section.

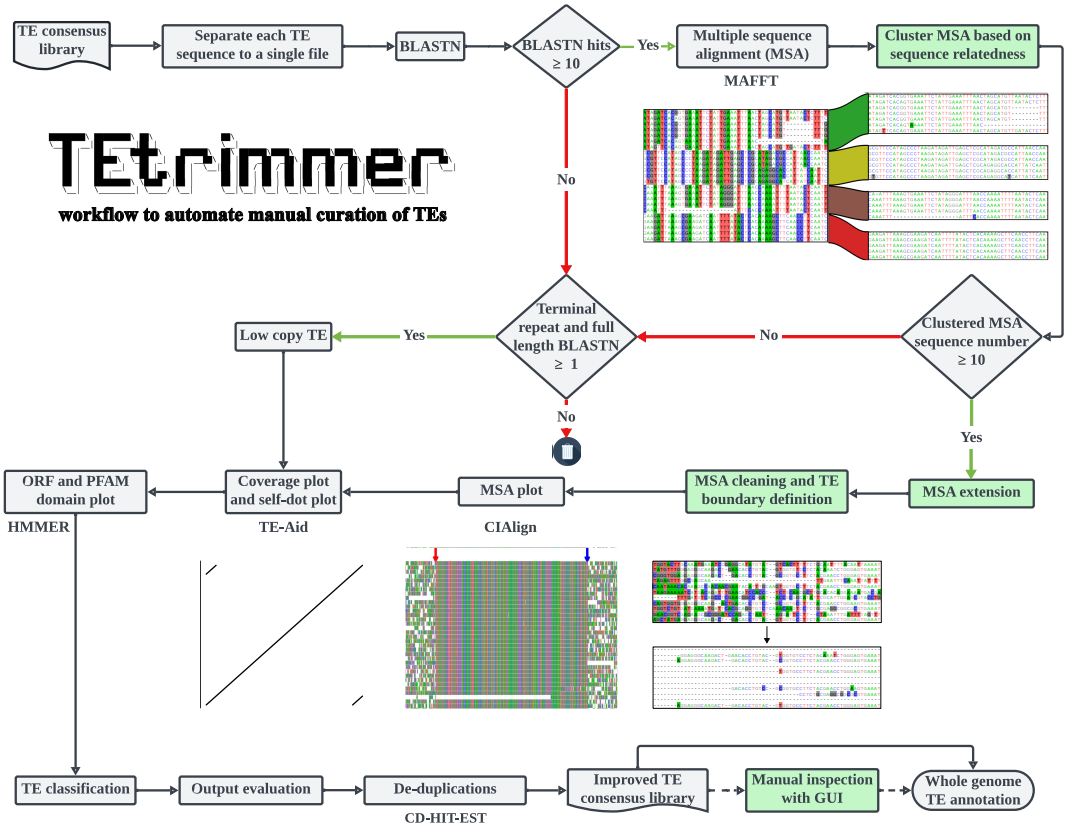


Figure 1: TEtrimmer workflow. TEtrimmer is a multi-thread tool to automate the manual curation of TEs. It can perform BLASTN searches of TE sequences retrieved from any TE consensus library against the respective genome and facilitate the generation of MSAs, MSA clustering, MSA sequence extension, MSA cleaning, TE boundary definition, and TE classification. Moreover, TEtrimmer provides detailed report plots and offers a graphical user interface (GUI) application to enable inspection and improvement of each output. The key features of TEtrimmer are highlighted by the green background color.

Table 1. Criteria for evaluation levels of the TEtrimmer output [a].

Category	Presence of terminal repeats	Classified	Sequence number in the MSA	Number of full-length BLASTN hits	Presence of PFAM domain
Perfect	TRUE	TRUE	≥30	≥5	TRUE
Good	TRUE	Not_required	≥10	≥2	Not_required
Reco_check	Not_required	Not_required	≥20	≥2	Not_required
Need_check	Not_required	Not_required	≥10	Not_required	Not_required
Low_copy	TRUE	Not_required	< 10	≥2	Not_required

[a] These criteria might not efficiently evaluate TEs lacking terminal repeats such as LINEs and SINEs.

TEtrimmer efficiently clusters sequences in the MSA

The TEtrimmer tool initially performs a BLASTN search for each input TE consensus sequence against the corresponding genome (Fig. 1). If more than ten hits are obtained, an MSA is established; otherwise the sequence is considered a potential low-copy TE. Due to the often high sequence similarity among

175 TE sub-families, this initial BLASTN search typically retrieves several types of TE sub-families per
176 input sequence. These can be visualized by distinct patterns of the aligned TE sequences corresponding
177 to the BLASTN hits (Fig. 2A). Thus, the first labor-intensive task of the traditional manual curation of
178 TEs is to separate TE sub-families within the MSA according to sequence relatedness. TEtrimmer
179 combines a phylogenetic tree approach (Fig. 2B) and a machine learning method (Density-Based Spatial
180 Clustering of Applications with Noise, DBSCAN) (Birant & Kut, 2007) to separate automatically
181 different alignment patterns that might correspond to distinct TE sub-families.

182 To illustrate the TEtrimmer clustering ability, one LTR retrotransposon from *Drosophila melanogaster*
183 identified by RepeatModeler2 was selected as an example. The BLASTN search of the selected TE
184 sequence against the *D. Melanogaster* genome yielded 99 hits. After extraction of the BLASTN hit
185 sequences and generation of an MSA, 101 divergent columns were selected in the original MSA and
186 combined into a new MSA of pseudo-sequences representing the divergent regions in the original MSA
187 (Fig. 2A, left part). Divergent columns are defined as columns where the proportion of the predominant
188 nucleotide in that column is below 0.8. The combined MSA based on the divergent columns was used
189 to construct an un-rooted maximum-likelihood phylogenetic tree (Fig. 2B), which was employed to
190 calculate a relative tree branch distance matrix (Supplementary Table 1). Finally, DBSCAN used the
191 distance matrix for MSA clustering, which yielded fourteen sub-MSAs ("clusters"), likely
192 corresponding to different TE subfamilies in the original MSA (Fig. 2A, right part and Fig. 2B).
193 Thirteen of these clusters, cluster 1 to 13, had clearly grouped sequences. The fourteenth cluster was
194 dubbed "noise", which means the sequences in this cluster could not be sub-grouped further. The
195 TEtrimmer MSA clustering efficiency was additionally supported by a principal component analysis
196 (PCA) plot based on the phylogenetic tree relative branch distance matrix (Fig. 2C). In the case of
197 clusters 1 to 13, the sequences in each cluster grouped adjacent to each other but were separated from
198 the sequences of other clusters. Regarding the "noise" cluster, its seven sequences were randomly
199 distributed across the entire PCA plot, indicating their low relatedness. This rather intricate example of
200 a complex MSA pattern illustrates that TEtrimmer can efficiently cluster TE sequences in an MSA.

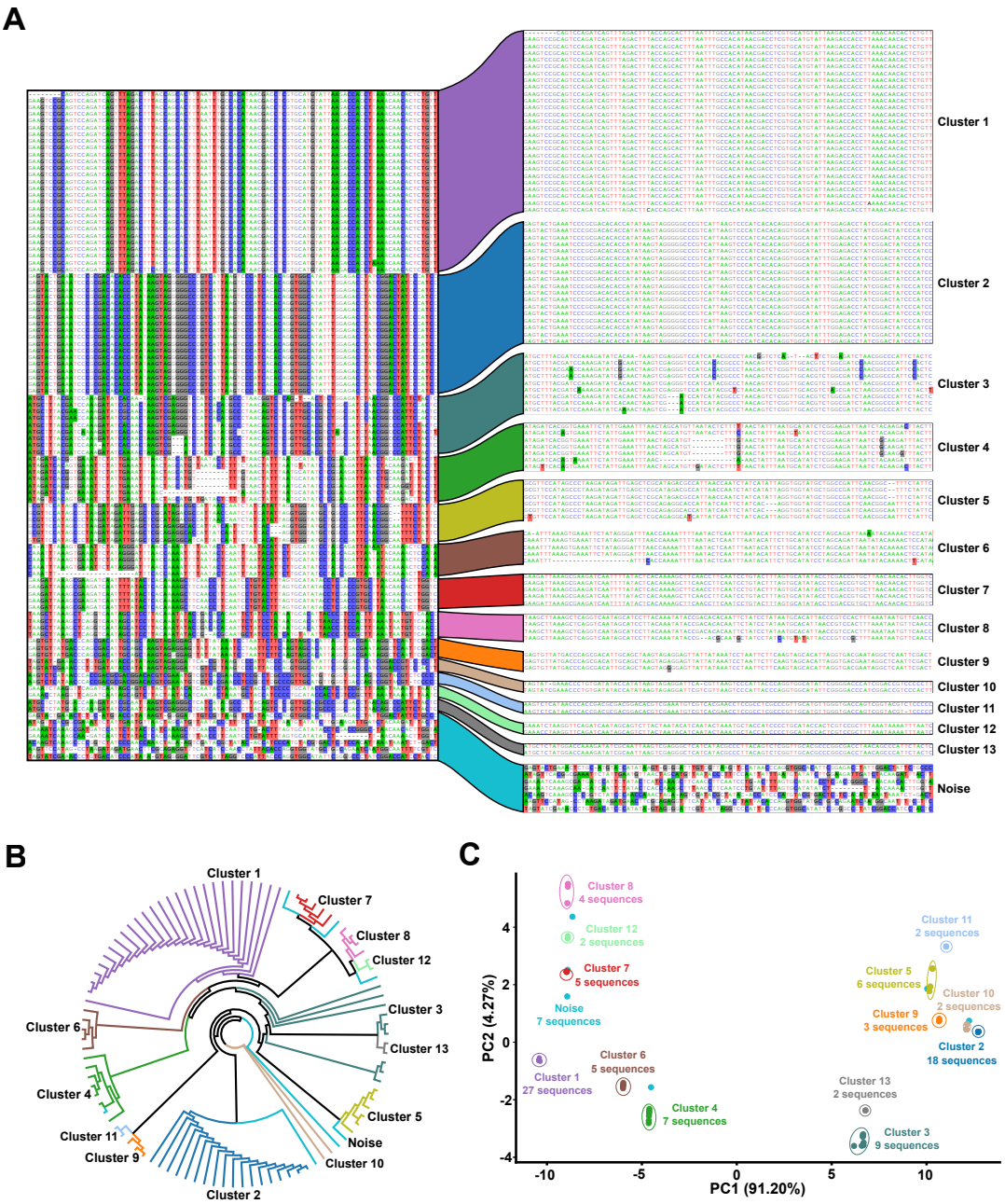


Figure 2: TEtrimmer efficiently clusters sequences within the MSA. One *D. melanogaster* LTR retrotransposon (named rnd-1-family-39), identified by RepeatModeler2, was chosen as an example for further analysis. TEtrimmer performed a BLASTN search of this LTR retrotransposon against the *D. melanogaster* genome, established an MSA with the respective BLASTN hit sequences, and conducted MSA clustering. **A** The left part represents the divergent columns in the MSA before clustering. The right area displays the MSAs after clustering by TEtrimmer based on the phylogenetic tree (B) relative branch distance-based DBSCAN analysis (epsilon = 0.1, min sample number = 2). Nucleotide background colors in the MSA represent sites where the proportion of the respective nucleotide in that column is below 0.4. The cluster names given here were consistently used in (B) and (C). Each color represents the corresponding cluster. **B** The un-rooted maximum likelihood phylogenetic tree generated by IQ-TREE is based on the MSA before clustering. Cluster labels and colors are consistent with (A). **C** PCA plot calculated using the phylogenetic tree relative branch distance matrix (B). The percentage of variance explained by the respective principal components PC1 and PC2 is indicated on the x- and

y-axis, respectively. Cluster labels and colors correspond with (A). The number of sequences within each cluster is indicated.

TEtrimmer efficiently extends and precisely cleans clustered MSA sequences

As elements retrieved by BLASTN searches can be incomplete, the sequences in the clustered MSAs typically need to be extended. TEtrimmer offers a convenient function for the expansion of sequence ends in the MSAs, which is iteratively used by the tool until the sequences cover the putative TE boundaries (Fig. 1; see Methods for further details on the sequence extension procedure). However, sequence extension beyond the actual TE borders usually results in non-TE sequences added to the MSA. These non-TE sequences, being typically very divergent, consistently produce badly aligned "noisy" regions in the MSA (Fig. 3A). Cleaning these regions is a time-consuming step of the manual curation process. The cleaning ("trimming") module of TEtrimmer efficiently addresses this task by removing gappy columns and the majority of lowly conserved regions within rows of MSAs after sequence extension. We demonstrate this capability by a representative MSA based on extended sequences derived from a *B. hordei* LINE element, rnd-1-family-34, (Fig. 3A). After MSA column cleaning by the TEtrimmer function "`remove_gap_columns`", the majority of gappy columns within the original MSA were eliminated (top and middle panel of Fig. 3A). Subsequently, the remaining noisy regions of the MSA were cleaned row by row by the TEtrimmer function "`crop_end_by_divergence`" (middle and bottom panel of Fig. 3A and B). Following column and row cleaning of the MSA, TE boundaries ("TE start" and "TE end") are determined by TEtrimmer (Fig. 3A and B, see Methods for details).

To evaluate the MSA cleaning performance, we compared manual cleaning to TEtrimmer-based cleaning for the above-mentioned LINE element. The effectiveness of cleaning conducted by TEtrimmer was assessed quantitatively using a customized confusion matrix (Provost et al., 1998), which showed a cleaning sensitivity of 0.977 and a precision of 0.999. This means that TEtrimmer correctly identified and removed 97.7% of the manually cleaned MSA regions and 99.9% of the regions it removed were in agreement with manual cleaning. An additional MSA row cleaning function of TEtrimmer not illustrated here is "`crop_end_by_gap`". In the TEtrimmer analysis process, this is an additional option preferentially used for LINE elements to deal in particular with their highly divergent 5′ regions. In contrast to the "`crop_end_by_divergence`" function, which is based on nucleotide proportions, "`crop_end_by_gap`" takes advantage of gap information for the cleaning process (see Methods for further details).

To test the cleaning performance across various types of TEs and different species, we randomly selected ten TEs from *B. hordei*, *D. melanogaster*, *D. rerio*, and *O. sativa*, including LTR retrotransposons, LINEs, and DNA transposons. After conducting BLASTN searches of the selected TE sequences against the corresponding genomes, sequence extension, MSA generation, and removal of gappy columns, we performed both manual and TEtrimmer-based MSA cleaning, the latter using the function "`crop_end_by_divergence`" with different thresholds. A detailed explanation of the cleaning thresholds can be found in the Methods section. TEtrimmer showed optimal row cleaning performance with a set threshold between 0.75 and 0.85, yielding averaged sensitivity and precision scores above 0.959 and 0.971, respectively (Fig. 3C). The results highlight the excellent MSA cleaning ability of TEtrimmer across various TE types from different species. Accordingly, the software name "TEtrimmer" relates to the advanced MSA trimming ability of the tool.

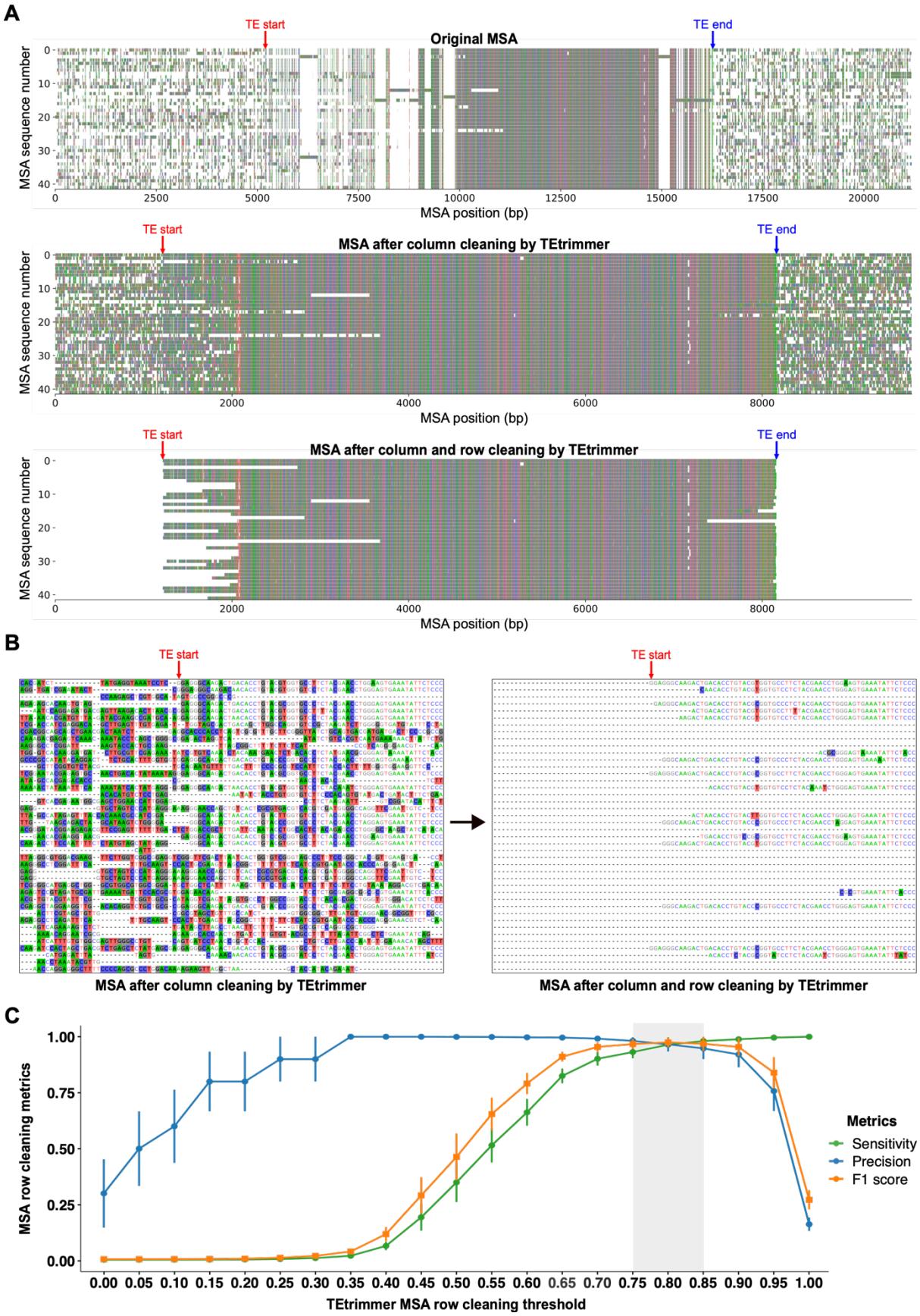


Figure 3: TEtrimmer succeeds in removing lowly conserved regions from MSAs. A *B. hordei* LINE element (named rnd-1-family-34), identified by RepeatModeler2, was chosen as a representative example for demonstrating the TEtrimmer MSA cleaning performance. **A** After BLASTN search of the

262 selected LINE sequence against the *B. hordei* genome, sequence extension, and MSA generation, the
263 LINE boundaries were identified, which are indicated by "TE start" (red) and "TE end" (blue).
264 Nucleotides are represented with colored bars; gaps are indicated as blank regions in the plots. The top
265 panel is the original MSA before cleaning, which contains many gappy columns and noisy rows. After
266 MSA column cleaning by the TEtrimmer function "`remove_gap_columns`", the majority of the
267 gappy columns are removed (middle panel). Then, TEtrimmer cleans sequences in the MSA row by
268 row using the TEtrimmer function "`crop_end_by_divergence`" (bottom panel), which removes
269 lowly conserved regions (bottom panel). **B** The left and right panels are both the magnified MSA regions
270 near the selected LINE element left boundary. The boundary is indicated by "TE start" (red). The left
271 panel illustrates the MSA before and the right panel after row cleaning by TEtrimmer via
272 "`crop_end_by_divergence`". Nucleotide background colors in the MSA represent sites where
273 the proportion of the respective nucleotide is below 0.4. **C** Ten TEs were randomly selected from the
274 RepeatModeler2 consensus libraries of *B. hordei* (barley powdery mildew fungus), *D. melanogaster*
275 (fruit fly), *D. rerio* (zebrafish), and *O. sativa* (rice). After BLASTN searches of the selected sequence
276 against the corresponding genomes, sequence extension, and MSA column cleaning, the generated
277 MSAs were used for benchmarking. Manual cleaning was performed to serve as a reference to enable
278 assessment of TEtrimmer MSA cleaning performance. The MSAs were cleaned by the TEtrimmer
279 cleaning function "`crop_end_by_divergence`" using cleaning thresholds ranging from 0 to 1 and
280 a sliding window size of 40 bp (a detailed description for thresholds and sliding windows can be found
281 in the Methods section). A confusion matrix analysis was conducted to evaluate the TEtrimmer
282 cleaning performance. The x-axis shows the cleaning threshold used by the TEtrimmer function
283 "`crop_end_by_divergence`"; the y-axis displays the confusion matrix score for the metrics
284 sensitivity (green), precision (blue), and F1 score (orange, calculated by (2 * (sensitivity * precision) /
285 (sensitivity + precision))). Standard error bars are shown in the plot. The grey-shaded box indicates the
286 threshold range where all three metrics scores are above 0.93.

287

288 **TEtrimmer provides detailed report plots and summary tables for each discovered TE**

289 Detailed report plots and summary tables for each output facilitate the assessment of outputs and
290 highlight differences in TE consensus sequences before and after TEtrimmer analysis (Fig. 4 and
291 Supplementary Table 2). A *B. hordei* LTR retrotransposon, named "ltr-1-family-22" and initially
292 identified by RepeatModeler2, was used for illustration of the report plots (Fig. 4). The first plot presents
293 the MSA columns around TEtrimmer-defined TE boundaries. The precise boundary positions are
294 denoted by colored arrows (Fig. 4A). This plot can help to inspect the existence of TSDs and the
295 accuracy of TE boundary definition. In the example, the marked boundaries clearly separate the highly
296 divergent from the conserved regions and TSD sequences can be identified (e.g. "GCACG" for the first
297 sequence in Fig. 4A), which represents a correct boundary definition as judged by manual inspection.
298 While the TE boundary plot only shows the ends of the MSA, the second plot displays the entire MSA
299 (Fig. 4B), which enables users to verify the alignment quality of the full MSA after TEtrimmer analysis.
300 The third type of plot illustrates the divergence of BLASTN hits against the TE consensus sequence
301 and their coordinates along the TE consensus sequence (Fig. 4C). For instance, the selected exemplary
302 LTR retrotransposon initially showed only one full-length BLASTN hit (Fig. 4C left panel), but 77 full-
303 length BLASTN hits were recovered after TEtrimmer analysis, which indicates a substantial
304 improvement (Fig. 4C right panel). The fourth type of plot is a self-dot plot, which can reveal TE
305 sequence features such as flanking LTRs and TIRs (Fig. 4D). The example self-dot plot after TEtrimmer
306 analysis shows flanking LTRs from bp 1 to 310 and bp 4638 to 4947 of the output consensus sequence
307 (Fig. 4D right panel), but only internal repeat regions (from bp 1167 to 1506 and bp 6003 to 6342),

308 indicative of a faulty TE, were identified for the input consensus sequence (Fig. 4D left panel). Besides
309 BLASTN hits and sequence features, ORF and protein domains are also crucial for evaluating a TE
310 consensus sequence. TEtrimmer provides a fifth type of plot to display the ORF and PFAM domains of
311 the consensus sequence before and after TEtrimmer analysis (Fig. 4E). For instance, after TEtrimmer
312 analysis ORF 2 was eliminated by deleting the sequence region from bp 1167 to 6002 of the input
313 consensus sequence. Additionally, ORF 1 and ORF 3 of the input consensus sequence (Fig. 4E, upper
314 panel) were fused into the single ORF 1 of the output consensus sequence (Fig. 4E, lower panel). The
315 final plot is a dot plot comparing TE consensus sequence similarity before and after TEtrimmer analysis
316 (Fig. 4F). The dot plot highlights both similar and different regions between the two sequences. For
317 example, the regions from bp 0 to 1166 and bp 6343 to 9508 of the input sequence were identical with
318 bp 311 to 1477 and bp 1478 to 4637, respectively, of the output sequence. In addition to the mentioned
319 plots, TEtrimmer provides a summary table compiling the analysis results such as the numbers of
320 BLASTN hits, sequence lengths, TE classifications, and the existence of terminal repeats
321 (Supplementary Table 2). The summary table helps to browse and index analysis results conveniently.
322 By combining the report plots and the summary table, users can quickly evaluate the accuracy of
323 TEtrimmer outputs and their differences from the input TE consensus sequence.

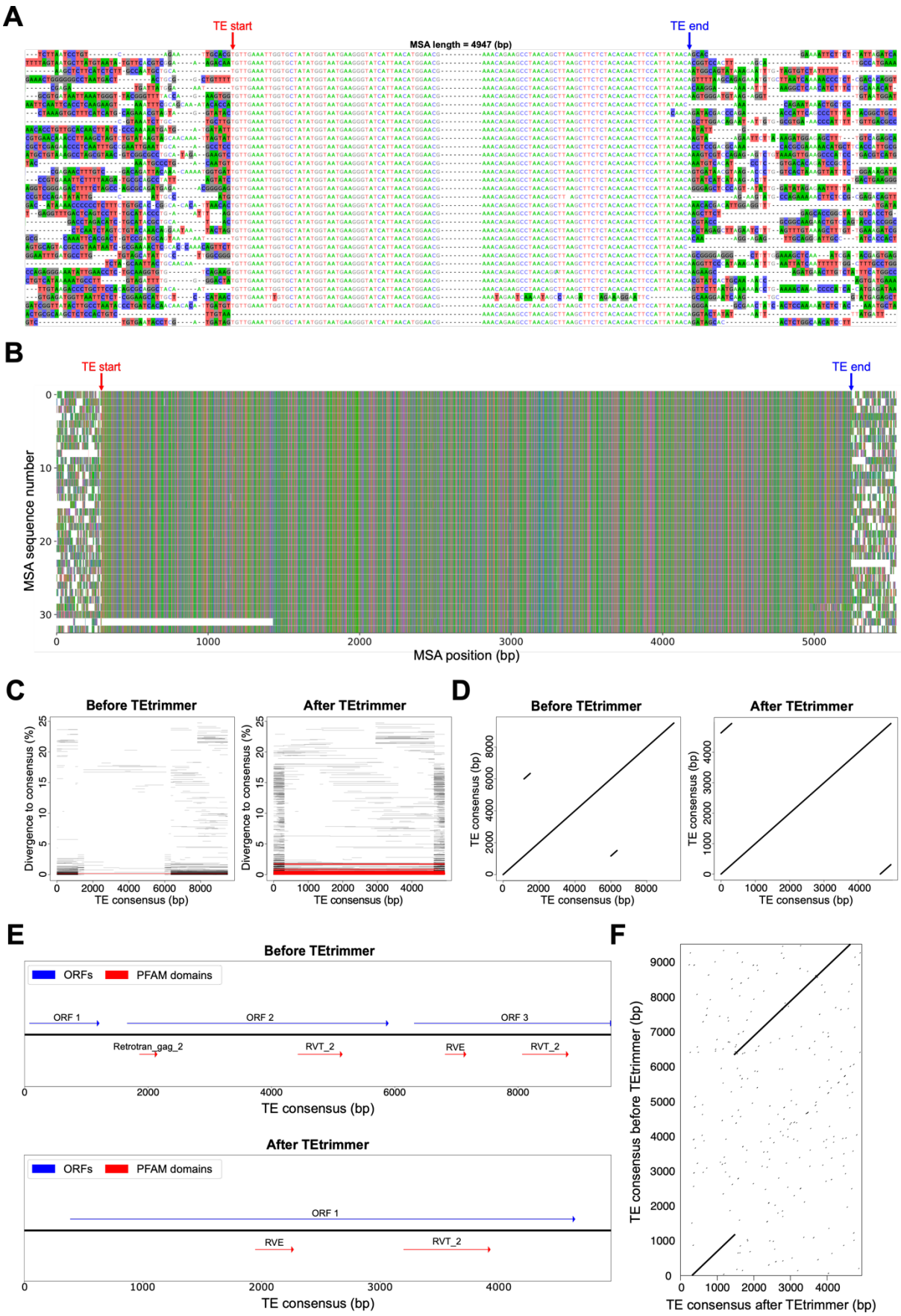


Figure 4: TEtrimmer provides comprehensive report plots for each discovered TE. The *B. hordei* LTR retrotransposon named "ltr-1-family-22", initially annotated by RepeatModeler2 and re-analyzed

with TEtrimmer, is shown as an example. **A** The alignment of 33 sequences in total shows the TE boundary regions of the MSA (100 nucleotide sites are displayed for each side) after TEtrimmer analysis. Nucleotide background colors in the MSA represent sites where the proportion of the respective nucleotide is below 0.4. The TE boundaries defined by TEtrimmer are indicated as "TE start" (red) and "TE end" (blue). The two boundary regions were artificially connected by ten gaps ("-") and the total length of the MSA is indicated on the top. **B** The plot shows the entire MSA after TEtrimmer analysis. The TE boundaries are indicated as in (A). Nucleotides are represented with colored bars; gaps are indicated as blank regions in the plot. The y-axis gives the nucleotide position (in bp) within the MSA. **C** The panels show a BLASTN plot before (left panel) and after (right panel) TEtrimmer analysis, each following a BLASTN search of the respective TE consensus sequence to identify TE sequence hits at a genome-wide scale. The x-axis indicates the TE consensus nucleotide position (in bp) and the y-axis is the sequence divergence in percent compared to the TE consensus sequence. Each line indicates a BLASTN hit; red lines highlight hits with a sequence divergence below 1.5% and a sequence coverage > 90%. **D** Self-dot plots before (left panel) and after (right panel) TEtrimmer analysis. The axes show the nucleotide position (in bp) in the TE consensus sequence. The repeat regions are represented by short diagonal lines outside the main diagonal. **E** The plot shows a genomic map of open reading frames (ORFs; blue arrows) and PFAM domains (red arrows) predicted in the TE consensus sequence before (upper panel) and after (lower panel) TEtrimmer analysis. The direction of the arrows indicates the strands of the respective ORFs/PFAMs in the sequence. ORF and PFAM identity are indicated on top of the arrows. After TE trimmer analysis, ORF2 is removed and ORF1 and ORF3 are fused to make the new ORF1. **F** Dot plots (window size = 25 bp, threshold = 50; the threshold represents the minimal sum of substitution scores in a defined window required for a dot to be plotted) of the TE consensus sequence before (y-axis) and after (x-axis) TEtrimmer analysis. Axes show the nucleotide position (in bp). Identical regions are represented by short diagonal lines outside the main diagonal.

TEtrimmer provides a user-friendly graphical user interface application for inspecting and improving discovered TEs

To examine and improve the de novo-generated TE consensus libraries established by TEtrimmer further, a GUI application is provided, which allows inspecting and correcting outputs conveniently. Four files are associated to each consensus sequence in the GUI: (1) the original MSA based on the BLASTN search of the input TE consensus sequence against the corresponding genome, (2) the MSA after clustering, sequence extension, and MSA cleaning but before TE boundary definition, (3) the MSA with defined TE boundaries used to generate the final TE consensus sequence, and (4) several report plots (Fig. 4). To group highly identical sequences, TEtrimmer calculates the nucleotide identity among all TE consensus output sequences. All four files mentioned above that relate to highly similar TE consensus sequences (identity >90%) are placed together into one "TE cluster" and listed by the GUI application (left window in Fig. 5).

For instance, after TEtrimmer analysis, the TE sequences "ltr_1_family_36", "rnd_1_family_139", and "rnd_1_family_187", which share more than 99.59% identity, and their associated files were placed into one "TE cluster" (Cluster 31; right window in Fig. 5). Moreover, multiple sequence dot plot files were provided to help visualizing consensus sequence similarities within a "TE cluster" (Supplementary Fig. 1). The multiple sequence dot plots help users conveniently inspect and compare similar outputs. Clustered files can be accessed by selecting the corresponding file names. Users can quickly assess each consensus sequence via the corresponding report plots file (Fig. 4) (e.g. "rnd_1_family_139#LTR__Copia_Perfect_4620_bp_99.59%.pdf"). If the TEtrimmer output appears to be inaccurate, users can easily modify and improve it using the provided MSA files and functions. For

example, if the sequence extension is insufficient in the view of the user, it can be manually elongated by activating the "Extend" button (right window in Fig. 5). The default extension size is 1000 bp for both ends, and these values can be easily modified separately for each side. After extension, a new file will appear with the extended sequences (e.g. "rnd_1_family_139#LTR__Copia.fa_1000_1000.fa", with 1000 bp extension on each side). Furthermore, to assist in identifying TE boundaries after manual extension, we integrated plot functions into the GUI application, allowing users to generate interactive report plots (Supplementary File 1) by activating the "TEAid" button. Additionally, the buttons "CropDiv", "CropGap", and "CleanCol" can be used to clean the MSA efficiently, which represent the same functions used for automated MSA cleaning in the TE trimmer pipeline (Fig. 3). In conclusion, the TEtrimmer GUI application integrates various files and functions associated with sequence extension, MSA cleaning, boundary identification, and visualization of TE consensus sequences, enabling users to examine and improve the TE consensus library conveniently. With the TEtrimmer GUI, the manual curation level of the TE consensus library can be readily achieved.

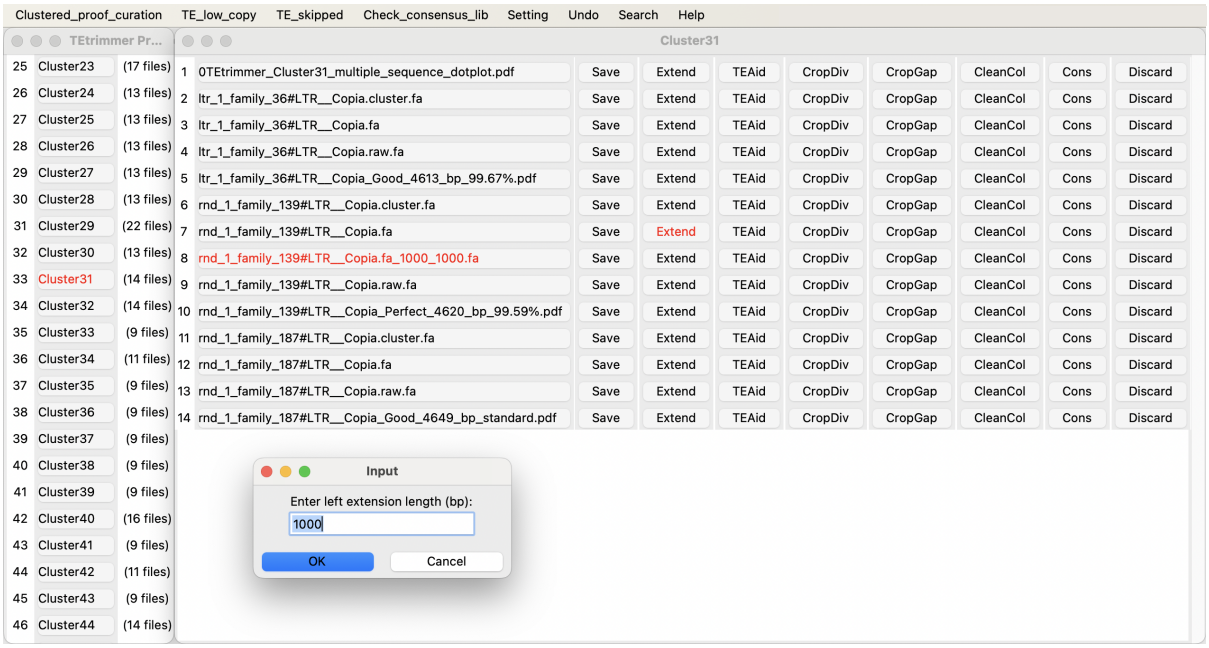


Figure 5: TEtrimmer provides a user-friendly graphical user interface (GUI) application to inspect and improve discovered TEs. Four files are associated with each consensus sequence (see main text for further details). All files are grouped together when their corresponding consensus sequences have >90% identity ("Cluster"). Users can inspect each result by selecting the corresponding file name. TE sequences in the MSA can be extended by activating the "Extend" button. The "TEAid" button helps to generate the interactive report plot (see Fig. 4). The buttons "CropDiv", "CropGap", and "CleanCol" relate to the MSA cleaning functions "`crop_end_by_divergence`", "`crop_end_by_gap`", and "`remove_gap_columns`", respectively. The button "Cons" can be used to generate a consensus sequence based on the corresponding MSA.

TEtrimmer improves the recovery of intact TEs compared to EDTA2 and RepeatModeler2

We tested the TEtrimmer performance on six organisms belonging to various kingdoms of eukaryotic life. These were *Blumeria hordei* (barley powdery mildew fungus), *Drosophila melanogaster* (fruit fly), *Danio rerio* (zebrafish), *Oryza sativa* (rice), *Zea mays* (maize), and *Homo sapiens* (human) (Table 2).

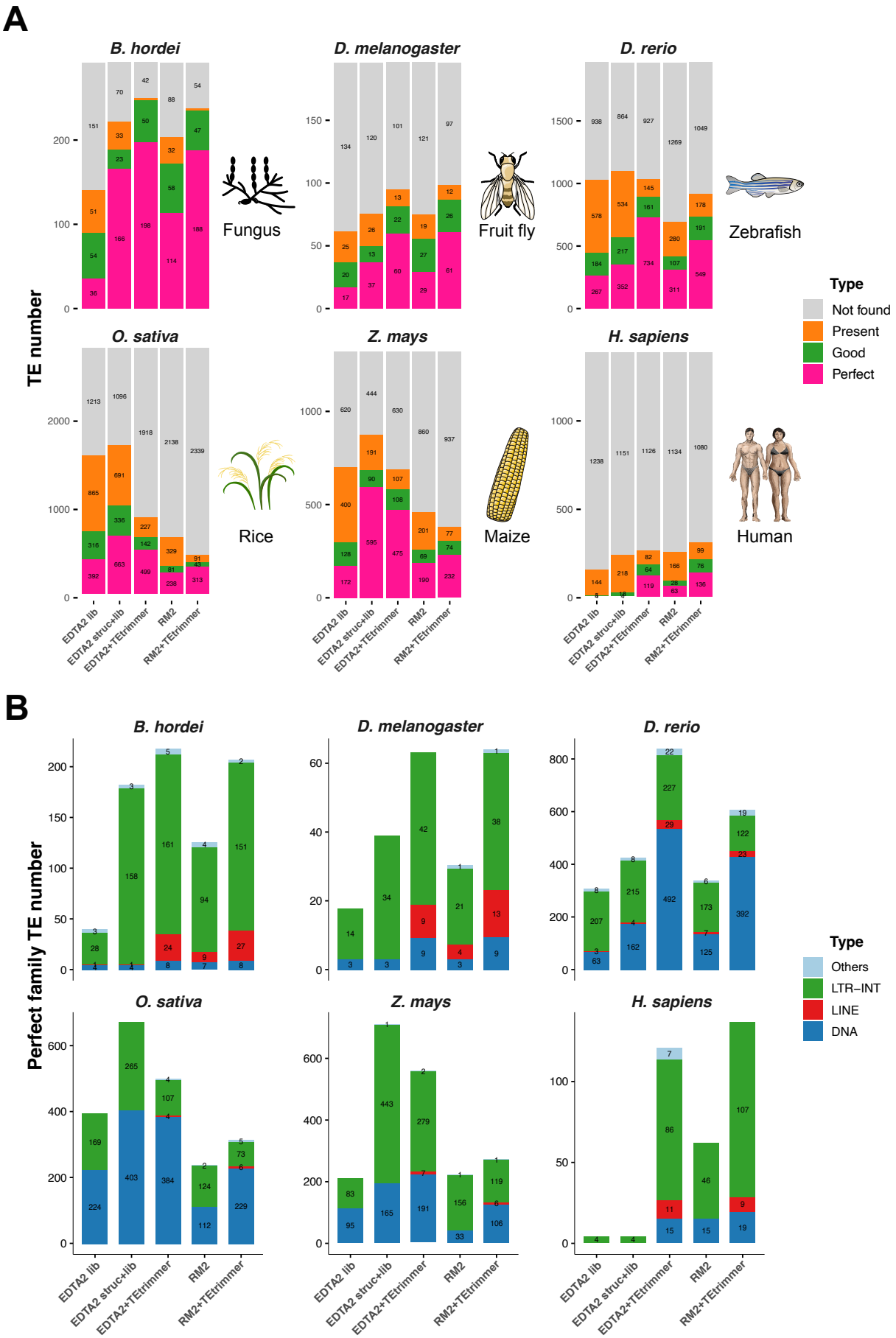
402 Benchmarking was initially conducted by directly comparing reference TE consensus libraries with de
403 novo-generated libraries derived from EDTA2, RepeatModeler2, and both these tools after additional
404 TEtrimmer analysis (consensus library after de-duplication). For this approach, we adopted the
405 benchmarking method reported by Flynn and co-workers (Flynn et al., 2020). Four benchmarking
406 levels were assigned to evaluate sequence matches between the reference libraries and the de novo-
407 generated libraries, including "Perfect", "Good", "Present", and "Not found". For the analysis, LTR
408 retrotransposons were separated into flanking LTRs and LTR internal sequences (LTR-INT) for all de
409 novo-generated and reference libraries. We found that up to 74% of the "Perfect" families were
410 comprised of flanking LTRs (Supplementary Table 3). To mitigate the potential bias caused by these
411 sequences, we eliminated all flanking LTRs for this benchmarking (see Methods for details).

412 **Table 2. Genomes and TE reference libraries used to benchmark TEtrimmer.**

Organism	TE reference library	Genome version	Genome link
Blumeria hordei	In-house manually curated	BGH_DH14_v4	https://www.ncbi.nlm.nih.gov/datasets/genome/GCA_900239735.1/
Danio rerio	Repbase	GRCz11	https://www.ncbi.nlm.nih.gov/datasets/genome/GCF_000002035.6/
Drosophila melanogaster	Repbase	Release 6 plus ISO1 MT	https://www.ncbi.nlm.nih.gov/datasets/taxonomy/7227/
Homo sapiens	Dfam	GRCh38.p14	https://www.ncbi.nlm.nih.gov/datasets/genome/GCF_000001405.40/
Oryza sativa	Repbase	version_7.0	http://rice.uga.edu/pub/data/Eukaryotic_Projects/o_sativa/annotation_dbs/pseudomolecules/version_7.0/all.dir/
Zea mays	Repbase	Zm-B73-REFERENCE-NAM-5.0	https://www.ncbi.nlm.nih.gov/datasets/genome/GCF_902167145.1/

413

414 The comparison between TEtrimmer results based on EDTA2 (EDTA2+TEtrimmer) and the EDTA2
415 consensus library (EDTA2 lib) revealed that TEtrimmer considerably improved the identification of
416 "Perfect" transposable element (TE) families by producing 1.3- to 29.8-times more "Perfect" TE
417 families across all six tested organisms (Fig. 6A). Specifically, EDTA2+TEtrimmer consistently
418 generated more DNA and LINE elements in the "Perfect" family category but fewer LTR
419 retrotransposons in case of *O. sativa* (Fig. 6B). EDTA2 struct+lib was generated by combining the
420 EDTA2 consensus library with all TE sequences discovered by the EDTA2 structure-based module,
421 followed by the removal of nearly identical sequences. EDTA2+TEtrimmer revealed more "Perfect"
422 families than EDTA2 struct+lib for most of the selected species, except for *O. sativa* and *Z. mays*.
423 TEtrimmer results based on RepeatModeler2 (RM2+TEtrimmer in Fig. 6A) generated 1.2- to 2.2-times
424 more "Perfect" families and consistently identified more DNA and LINE elements as "Perfect" families
425 across all selected organisms compared to RepeatModeler2 alone (RM2 in Fig. 6A). Conversely, fewer
426 LTR retrotransposons were identified by TEtrimmer for *D. rerio*, *O. sativa*, and *Z. mays* (Fig. 6B and
427 Supplementary Table 4), which might be due to the relatively inefficient identification of low-copy TEs
428 by TEtrimmer. Overall, TEtrimmer predominantly improved the curation of TEs in all organisms except
429 for LTR retrotransposons in the two plant genomes.


A
B. hordei
D. melanogaster
D. rerio
O. sativa
Z. mays
H. sapiens
Fungus
Fruit fly
Zebrafish
Rice
Maize
Human
TE number
Type
Not found
Present
Good
Perfect
EDTA2 lib
EDTA2 struc+lib
EDTA2+TEtrimmer
RM2
RM2+TEtrimmer
B
Perfect family TE number
Others
LTR–INT
LINE
DNA

430

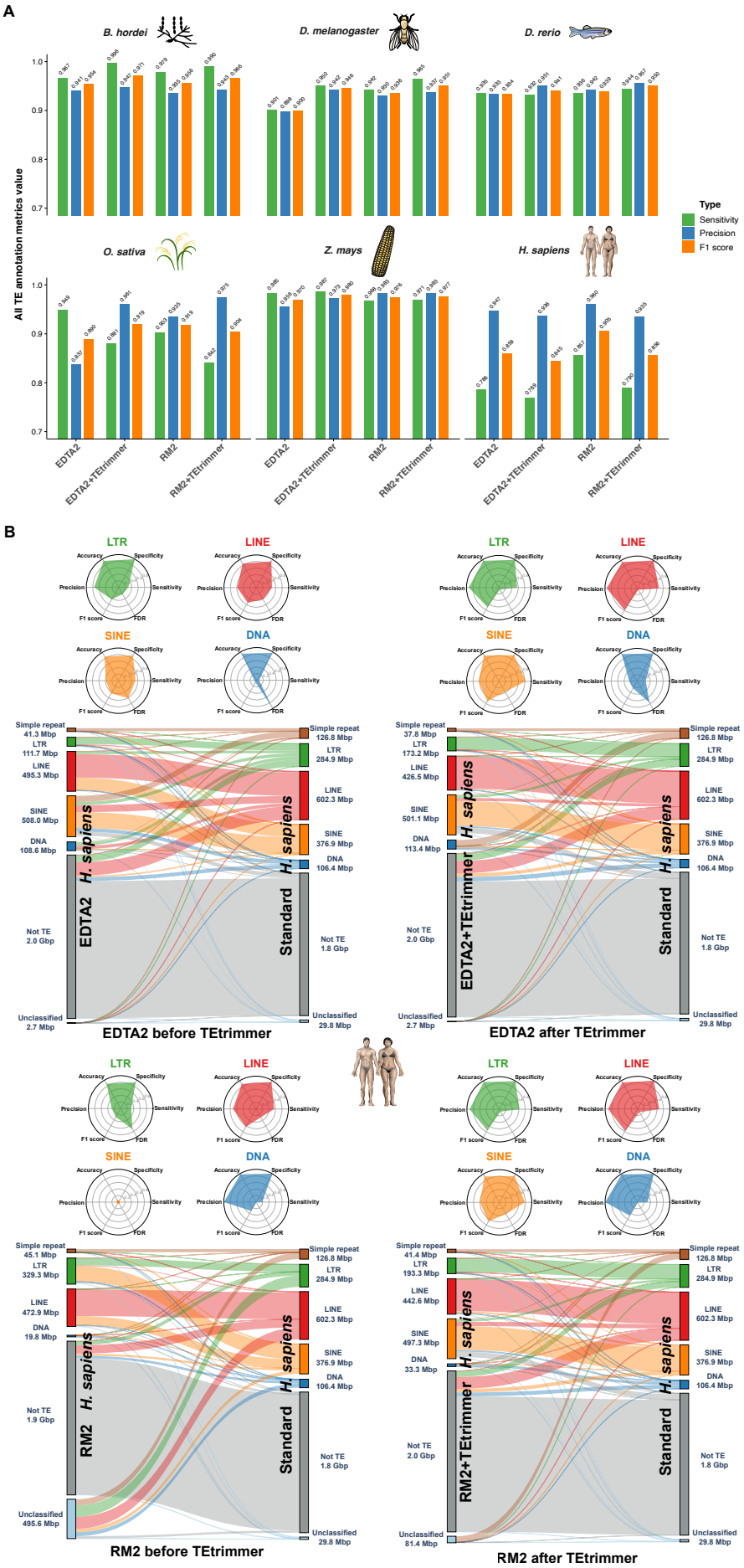
Figure 6. TEtrimmer improves the discovery of intact TEs compared to EDTA2 and RepeatModeler2. We benchmarked the performance of EDTA2, RepeatModeler2 (RM2), and both these tools after additional TEtrimmer analysis (EDTA2+TEtrimmer and RM2+TEtrimmer, respectively) by comparing the TE quality in the de novo-generated consensus libraries with the quality in the corresponding reference TE consensus libraries (Table 2) of the genomes of six organisms, i.e., *B. hordei* (barley powdery mildew fungus), *D. melanogaster* (fruit fly), *D. rerio* (zebrafish), *O. sativa* (rice), *Z. mays* (maize), and *H. sapiens* (human). Two libraries were used for EDTA2, termed "EDTA2 lib" and "EDTA2 struct+lib". "EDTA2 lib" represents the EDTA2 consensus library and "EDTA2 struct+lib" was generated by combining the EDTA2 consensus library with all TE sequences discovered by the EDTA2 structure-based module. **A** The stacked bar graphs show the proportion of correctly discovered consensus TEs by the respective analysis tools/pipelines indicated on the x-axis according to the color code shown on the right. The y-axis displays the number of TE sequences in the respective reference TE library, i.e., 292 for *B. hordei*, 196 for *D. melanogaster*, 1967 for *D. rerio*, 2786 for *O. sativa*, 1320 for *Z. mays*, and 1391 for *H. sapiens*. Benchmarking categories included "Perfect" (pink), "Good" (green), "Present" (orange), and "Not found" (grey) (Flynn et al., 2020). "Perfect" means that the reference TE consensus sequences have one match in the de novo-generated library with > 95% similarity and coverage. "Good" indicates that the reference TE consensus sequence has multiple overlapping matches in the de novo-generated library, each with > 95% similarity and full coverage. "Present" is similar to "Good", but the required minimum similarity and total coverage is decreased to 80%. The remaining TE sequences were assigned to the "Not found" category. **B** The stacked bar graphs show the number of "Perfect" matches in (A) resolved for the TE types of DNA transposons, LINEs LTR retrotransposon internal sequences (LTR-INT), and others (including e.g. Helitrons, SINEs, and simple repeat elements). The x-axis displays the tools/pipelines and the y-axis the number of TE sequences.

TEtrimmer exhibits superior genome-wide TE annotation ability in comparison to EDTA2 and RepeatModeler2

In addition to the direct comparison of TE consensus libraries, we benchmarked EDTA2, RepeatModeler2, and TEtrimmer concerning whole-genome TE annotation by using a confusion matrix (Ou et al., 2019; Provost et al., 1998). First, all genome regions annotated as any type of TE, termed "all TEs", were analyzed to assess the accuracy of TE masking in the genome. TEtrimmer exhibited higher sensitivity and precision (F1 scores > 0.941) than EDTA2 and RepeatModeler2 for *B. hordei*, *D. melanogaster*, *D. rerio*, and *Z. mays*, except for a slightly lower sensitivity score for the *D. rerio* EDTA2+TEtrimmer library compared to EDTA2 alone (Fig. 7A). By contrast, for *O. sativa* (rice), TEtrimmer exhibited lower sensitivity but higher precision than EDTA2 and RepeatModeler2. TEtrimmer also showed lower sensitivity and precision for *H. sapiens* compared to the other tools (Fig. 7A and Supplementary Table 5).

Due to the overall poor performance on TE consensus library construction for *H. sapiens* by all tested tools (Fig. 6A) and TEtrimmer's relatively lower TE annotation score for *H. sapiens* (Fig. 7A), we calculated a confusion matrix for different TE types (radar plots in Fig. 6B). In case of LTR retrotransposons, LINEs, SINEs, and DNA transposons, both the sensitivity and precision were considerably improved by additional TEtrimmer analysis. Moreover, genome-wide TE annotation based on the TEtrimmer consensus libraries exhibited better agreement with the reference annotation than those from EDTA2 and RepeatModeler2 alone (Sankey plots in Fig. 7B). For instance, the TE annotation from RepeatModeler2 (before TEtrimmer) yielded 495.6 Mbp of unclassified TEs and no SINE elements for *H. sapiens* (Fig. 7B lower panel), while additional TEtrimmer analysis reduced the

477 unclassified TE category to 81.4 Mbp and annotated 497.3 Mbp of SINEs with high accuracy, (Fig. 7B).
478 In conclusion, TEtrimmer libraries generally outperformed EDTA2 and RepeatModeler2-derived
479 libraries for whole-genome TE annotation across most of the tested organisms. Additionally, TEtrimmer
480 provided more correct TE classifications.


A
B. hordei
D. melanogaster
D. rerio
O. sativa
Z. mays
H. sapiens
All TE annotation metrics value
Type
Sensitivity
Precision
F1 score
EDTA2
EDTA2+TEtrimmer
RM2
RM2+TEtrimmer
B
LTR
LINE
SINE
DNA
Accuracy
Specificity
Sensitivity
FDR
F1 score
Precision
H. sapiens
Standard
EDTA2 before TEtrimmer
EDTA2 after TEtrimmer
RM2 before TEtrimmer
RM2 after TEtrimmer

481

482 **Figure 7: TEtrimmer exhibits superior genome-wide TE annotation ability in comparison to**
483 **EDTA2 and RepeatModeler2.** We assessed the genome TE annotation performance of EDTA2,
484 RepeatModeler2 (RM2), and both after additional TEtrimmer analysis (EDTA2+TEtrimmer and
485 RM2+TEtrimmer, respectively). All genome-wide TE annotation results based on de novo-generated
486 libraries were compared with the results of the respective TE reference libraries (Table 2). We used a
487 confusion matrix (Ou et al., 2019; Provost et al., 1998) to calculate the sensitivity, precision, accuracy,
488 specificity, F1 score, and false discovery rate (FDR) of the tools. The genomes of six organisms were
489 used for the analysis, i.e., *B. hordei* (barley powdery mildew fungus), *D. melanogaster* (fruit fly), *D.*
490 *rerio* (zebrafish), *O. sativa* (rice), *Z. mays* (maize), and *H. sapiens* (human). **A** The bar plots show the
491 sensitivity (green), precision (blue), and F1 score (orange) calculated with a confusion matrix for the
492 six organisms indicated on top of each graph. The x-axis represents the respective analysis tools, and
493 the y-axis displays the metrics score values based on overall genome TE annotation correctness ("All
494 TEs") at a range of 0.7 to 1.0. **B** Detailed genome-wide TE annotation benchmarking for *H. sapiens*.
495 The radar plots in the upper panels show the benchmarking metrics for LTR retrotransposon (green),
496 LINEs (red), and SINEs (orange). The lower panels display the annotation overlap and differences
497 between de novo-generated TE libraries from the indicated tool (left) and the reference consensus TE
498 library for *H. sapiens* (right); each bar indicates the number of Mbp masked as the respective element
499 in the genome. The links between the left and right bars indicate the connections between the respective
500 libraries. TE types shown are LTR retrotransposon (green), LINE (red), SINE (orange), DNA
501 transposon (dark blue), unclassified (light blue), and genomic sequence not identified as TE ("Not TE";
502 grey).

503

Discussion

Due to decades of genome sequencing and the advanced long-read third-generation sequencing technology, the number of high-quality genomes is growing rapidly. However, precise and accurate TE annotation in genomes remains a major challenge. Several tools have been developed to automate TE discovery and annotation. Nonetheless, the manual curation of TEs is still mandatory to obtain comprehensive and detailed insights into the TE landscape of a given genome. Manual curation is a time-consuming and challenging process and often beyond the scope of a given research project. In this study, we introduce TEtrimmer, a novel software designed to automate and support the manual curation of TEs. TEtrimmer efficiently clusters and cleans MSAs and precisely defines TE boundaries. Moreover, TEtrimmer provides comprehensive report plots and a user-friendly GUI application to facilitate the inspection and improvement of discovered TE consensus sequences. In particular, the sliding window-based cleaning strategy of the MSA, which results in superior precision, and the interactive GUI application are innovative features of TEtrimmer as compared to other related software tools.

In manual curation procedures, the first labor-intensive step is separating the MSA into different clusters according to the TE sequence similarity (Fig. 2). To accelerate this step, we combined the selection of distinct MSA columns, a phylogenetic tree approach, and a machine learning method (DBSCAN) to cluster MSAs automatically and precisely. Although other clustering tools exist, none of these are fully automated. For example, TreeCluster can group phylogenetic trees according to the relative branch distance (Balaban et al., 2019), but the final cluster number must be defined manually. Furthermore, CD-HIT-EST can cluster TE sequences without generating an MSA (Fu et al., 2012), but the identity threshold needs to be manually adjusted for each MSA separately. MCHelper (Orozco-Arias et al., 2023) also provides an MSA clustering function and, like TEtrimmer, relies on DBSCAN. However, MCHelper does not include the selection of divergent MSA columns and lacks the phylogenetic tree approach. It instead uses the K2P (Kimura 2-Parameter) sequence distance matrix for DBSCAN analysis. Selectively using divergent MSA columns for clustering is crucial because consistent columns (i.e., regions of high conservation) can potentially mask or dilute the differences among sequences. Furthermore, in our experience, an un-rooted maximum likelihood phylogenetic tree is superior to calculate sequence distances for processing by DBSCAN. In contrast to the tools mentioned above, the TEtrimmer clustering algorithm automatically defines the appropriate cluster number and thresholds. It combines the selection of divergent MSA columns and a phylogenetic tree strategy for a more advanced DBSCAN clustering analysis. This approach is not only useful for the clustering of TE sequences but also for any set of nucleotide or amino acid sequences. However, because the clustering module of TEtrimmer relies on MSAs, it cannot handle large-scale datasets containing thousands of DNA sequences.

Besides MSA clustering, we developed a new MSA cleaning algorithm to remove highly divergent regions in MSAs in a specific manner (Fig. 3). Removing divergent sites from MSAs is important because they can significantly hinder the generation of TE consensus sequences and the definition of TE boundaries. Fragmented TEs and excessive end extension during the curation process typically result in the addition of many TE-unrelated flanking sequences in the MSA. This issue is especially pronounced in the case of the 5'-ends of LINE elements, which tend to be highly degraded (Goubert et al., 2022) (Fig. 3). Other tools such as trimAI (Capella-Gutiérrez et al., 2009), Clipkit (Steenwyk et al., 2020) and CIAlign (Tumescheit et al., 2022) are able to perform MSA cleaning but are mainly designed to remove gappy columns and optimize MSAs for better phylogenetic tree construction. CIAlign provides basic functions to clean rows and columns of MSAs but mainly relies on alignment gap information, which is insufficient for TE-related MSA scenarios. TEtrimmer introduces a sliding

window strategy to analyze sequentially the entire MSA nucleotide by nucleotide. Our benchmarking of the cleaning performance and accuracy (Fig. 3C) demonstrates that the TEtrimmer MSA cleaning module exhibits a performance close to that of manual MSA cleaning. In summary, TEtrimmer is to the best of our knowledge the first tool that provides extensive automated MSA cleaning procedures for TE sequences.

TEtrimmer provides a powerful GUI application to facilitate the inspection and improvement of its outputs. This can close the gap between de novo-based TE libraries and the traditional, manual curation-based TE consensus libraries. TEtrimmer is the first tool to introduce a TE discovery- and curation-related GUI application. Moreover, we offer more TE-related functions in this GUI application. For example, users can not only examine and improve TEtrimmer outputs but also any TE consensus libraries established by other tools such as EDTA2 and RepeatModeler2 (Supplementary Fig. 2). They can further easily perform BLASTN searches, MSA generation, MSA sequence extension, MSA cleaning, MSA plotting, TE-Aid style plotting, and consensus sequence generation. The integration of TE benchmarking functions into the GUI application is a future option.

To test the performance of TEtrimmer regarding the discovery of intact TEs, we executed TEtrimmer using the outputs from EDTA2 and RepeatModeler2, respectively, based on six organisms of different kingdoms of eukaryotic life. The TE consensus libraries generated by TEtrimmer consistently contained a higher number of intact ("Perfect") TEs than the consensus libraries generated by EDTA2 and RepeatModeler2 (Fig. 6A). By contrast, TEtrimmer discovered less intact LTR retrotransposons for *D. rerio*, *O. sativa* and *Z. mays* in some cases. A possible explanation for this contrasting performance could be that TEtrimmer mainly relies on the repetitive nature of TEs to improve the consensus library and, in turn, may not be effective in discovering accurately low-copy TEs. In contrast to TEtrimmer, both EDTA2 and RepeatModeler2 rely on structure-based methods to identify low-copy LTR retrotransposons. Nevertheless, such low-copy TEs can be easily recovered by TEtrimmer with the provided GUI application through the optional review and curation process.

In addition to the comparative assessment of TE integrity, we introduced a confusion matrix (Ou et al., 2019) to evaluate the TE annotation performance at a genome-wide scale. EDTA2 and RepeatModeler2 exhibited great sensitivity and precision for the selected organisms at the level of "all TEs", which indicates that these tools can accurately mask TE regions in the respective genomes. However, most of the benchmarking metrics were improved after TEtrimmer analysis, except for *O. sativa* and *H. sapiens* (Fig. 7A) where the metrics values were slightly decreased, most likely due to the relatively poor performance of TEtrimmer regarding the analysis of low-copy TEs. Nevertheless, we found an overall improved TE classification accuracy for *H. sapiens* by TEtrimmer (Fig. 7B). Here, we introduced for the first time Sankey plots to the TE field to visualize effectively the quality of TE annotation compared to the respective reference library.

To date, EarlGrey and MCHelper are the two most prominent tools designed for automating the manual curation of TEs. TEtrimmer outperforms these tools in five main aspects. (1) EarlGrey does not provide a clustering function for MSAs, which can be a limitation for analyzing highly divergent TEs. Compared to MCHelper, TEtrimmer has a more sophisticated MSA clustering strategy that is based on the selection of divergent MSA columns in combination with the generation of a phylogenetic tree, which can dramatically improve DBSCAN performance. (2) Both EarlGrey and MCHelper are not able to clean efficiently MSAs, due to a lack of functions to clean MSA rows. (3) For each iteration of MSA extension, EarlGrey and MCHelper extend all TE sequences in the MSA. By contrast, we developed an MSA extension algorithm that can selectively extend TE sequences based on the divergence in the alignment. This strategy facilitates even the analysis of highly degraded regions, such as the 5' regions of LINE elements. (4) TEtrimmer assigns different evaluation levels for each output based on criteria

such as the presence of terminal repeats, classification status, MSA sequence number, number of full-length BLASTN hits, and PFAM domain prediction (Table 1). This helps in systematically categorizing and assessing the quality of the outputs. (5) TEtrimmer provides comprehensive report plots (Fig. 4) and a user-friendly GUI application (Fig. 5), which allow users to inspect and improve each TEtrimmer output sequence conveniently. We also integrated the MSA cleaning, MSA extension, and plotting functions into the GUI application to achieve manual curation-level TE libraries efficiently.

Conclusions

In conclusion, TEtrimmer is a powerful and user-friendly tool designed to automate the manual curation of TEs, addressing the major challenge of accurately annotating these repetitive DNA sequences in eukaryotic genomes. By integrating advanced techniques for MSA clustering, cleaning, and extension, as well as TE boundary definition, TEtrimmer enhances the quality of TE annotations markedly. The tool provides detailed report plots and a GUI application for efficient inspection and refinement of results, making it accessible even for researchers that lack extensive levels of expertise in TE genetics. Comprehensive benchmarking against the genomes of six diverse eukaryotic organisms demonstrated the advanced performance of TEtrimmer compared to existing tools like RepeatModeler2 and EDTA2, particularly in identifying intact (full-length) TEs. Hence, TEtrimmer bridges the gap between automated TE annotation and the gold standard of manual curation, offering a robust solution for the accurate and efficient annotation of TEs in genomic studies.

Methods

BLASTN searches

TEtrimmer (v1.3.0) uses the TE consensus library output from any de novo TE discovery tool such as EDTA2, RepeatModeler2, and REPET, or consensus TE libraries from closely related species, as input to improve the TE library quality. First, TEtrimmer removes duplicated input sequences via CD-HIT-EST (Fu et al., 2012) (v.4.8.1) when the "`--dedup`" option is enabled. Sequences with more than 95% sequence identity and coverage are merged into one element. Then, each input sequence is converted into a single file to enable multi-thread analysis. TEtrimmer uses these separated sequences to perform BLASTN searches against the respective genome to find query copies for each TE sequence. By default, only the 70 longest and 30 randomly selected BLASTN hits are used for further analysis when more than 100 target sequences are identified by the BLASTN query, as these are typically sufficient for meaningful downstream analysis. Finally, the BLASTN search result is converted into a BED file.

MSA generation

TE sequences are extracted based on the BED file derived from the BLASTN search of each input TE consensus sequence against the corresponding genome (see above) by "`bedtools getfasta`" (Quinlan & Hall, 2010) (v2.31.1). Those sequences are aligned by MAFFT (Katoh & Standley, 2013) (v7.520) with the fast alignment option "`--retree 1`" and any gappy alignment columns are removed prior to MSA clustering by the function "`remove_gap_columns`" (see section "MSA cleaning" below).

MSA clustering

Because of the typically low sequence divergence among TE subfamilies, several types of sub-families are usually included in the MSA based on the BLASTN search derived from each input consensus sequence against the respective genome (Fig. 2). To separate these subfamilies, TEtrimmer calculates the percentage of each type of nucleotide in every column. Gaps in that column are not included in the calculation. Divergent columns are defined as those where all nucleotide proportions are lower 0.8 in that column. All divergent columns are merged together into a new FASTA file, yielding an MSA of pseudo-sequences representing the divergent regions in the original MSA, which is used by IQ-TREE (Minh et al., 2020) to generate a maximum likelihood phylogenetic tree. We use the fixed IQ-TREE model "K2P+I" to reduce the execution time of this module. Then, a sequence distance matrix is calculated based on the relative tree branch distances. Finally, the DBSCAN (Birant & Kut, 2007) machine learning method is performed based on the sequence distance matrix to cluster the sequences of the MSA, with epsilon set to 0.1 and minimum samples set to 2. More than two clusters can be generated after MSA clustering. By default, TEtrimmer only proceeds with the top two clusters containing the highest number of sequences, which provides in our hands typically a good balance between sensitivity and computational intensity. Users can increase this number by the option "`--max_cluster_num`" but need to consider that increasing the cluster number can cause a high rate of false-positive results (i.e., non-TE sequences mistakenly classified as TEs).

MSA cleaning ("trimming")

For cleaning of columns within an MSA, TEtrimmer uses the function "`remove_gap_columns`". By default, TEtrimmer removes columns when the gap comprises more than 80% of the column or the nucleotide number in this column is less than five. In a subsequent step, columns with a gap percentage between 40% and 80% and where the predominant nucleotide constitutes less than 70% of the total nucleotides are deleted.

658 In addition to gappy columns, highly divergent sequence regions might occur in the MSA. TEtrimmer
659 includes two functions to remove divergent regions from the ends of the MSA row by row. The function
660 "`crop_end_by_divergence`" first calculates the proportion of each nucleotide in every column
661 that have at least five nucleotides. Gaps are not included in this calculation. All nucleotide proportions
662 in columns with fewer than five nucleotides are considered as zero. Then the entire MSA is converted
663 into a proportion matrix by replacing each nucleotide with the corresponding proportion value and all
664 gap positions are converted to zero. For each row of the proportion matrix, two sliding windows (default
665 size 40 bp) are created at both ends that move stepwise towards the opposite end of the sequence
666 nucleotide position by nucleotide position. Each sliding window stops when the mean of the nucleotide
667 proportions within the sliding window is greater than the set threshold (default 0.7), indicative of having
668 reached a conserved region. For each sliding window starting from the left end of the sequence, all
669 nucleotide positions between the left end of the sequence and the stopping point (left boundary) of this
670 sliding window are deleted. Similarly, for each sliding window that starts from the right end of the
671 sequence, all nucleotide positions between the right end of the sequence and the stopping point (right
672 boundary) of the sliding window are deleted. After completion of all rows, the process is repeated with
673 a smaller sliding window (default size 4 bp) and a higher threshold (default 1.0) for fine-polishing the
674 MSA ends.

675 The second function used to clean the MSA is called "`crop_end_by_gap`". It deploys a similar
676 sliding window cleaning principle as "`crop_end_by_divergence`", but the difference is that
677 "`crop_end_by_gap`" will not convert the MSA into a proportion matrix. Instead, it takes advantage
678 of the original MSA. For each sequence in the MSA, two sliding windows (default size 250 bp) are
679 created at both ends that stepwise move towards the opposite end of the sequence nucleotide position
680 by nucleotide position. Each sliding window stops when the proportion of gaps included in the sliding
681 window is below the set threshold (default 0.1), indicative of having reached a conserved region. For
682 each sliding window that starts from the left end of the sequence, all nucleotide positions between the
683 left end of the sequence and the stopping point (left boundary) of this sliding window are deleted.
684 Similarly, for each sliding window that starts from the right end of the sequence, all nucleotide positions
685 between the right end of the sequence and the stopping point (right boundary) of the sliding windows
686 are deleted. The "`crop_end_by_gap`" function is also used to facilitate MSA extensions (see below).

687 **MSA sequence extension**

688 Annotated TEs derived from de novo TE discovery software can be fragmented and truncated. To
689 complete such TE sequences, TEtrimmer iteratively extends both ends of the sequences from the
690 clustered MSA separately via "`bedtools slop`" (Quinlan & Hall, 2010). The default step size is
691 1000 bp for each end per extension, with a maximum total extension size of 7000 bp at each end of the
692 sequences.

693 First, TEtrimmer performs left end sequence extension of the MSA. To save computing time, after each
694 round of sequence extension TEtrimmer only aligns the newly extended sequences from the current
695 round, followed by cleaning of gappy columns via the function "`remove_gap_columns`" as
696 described above (see section "MSA cleaning"). Then, the function "`crop_end_by_gap`" (see section
697 "MSA cleaning" above) is employed to clean the MSA additionally row by row, and the sum of the
698 removed sites for each sequence (row) is recorded. The sequence is excluded from the next round of
699 extension when the sum of its removed sites by "`crop_end_by_gap`" is more than 90% of the
700 sequence length, indicating a sufficient extension for this sequence as seemingly flanking genomic
701 regions have been reached. Moreover, to assess if the overall left end extension is sufficient to include
702 the left TE boundary, the newly extended and cleaned MSA (only containing the sequences extended

703 in the current round) is used to generate a consensus sequence where the letter "N" represents any
704 ambiguous position. An ambiguous position is an MSA column site where the proportion of the
705 predominant nucleotide is below 0.7 or the total nucleotide count is below five. Then, a sliding window
706 (default size 150 bp) is created at the left end of the TE consensus sequence that moves stepwise towards
707 the opposite end of the sequence nucleotide position by nucleotide position. The sliding window stops
708 when the proportion of the ambiguous letter "N" (as defined above) within the sliding window is less
709 than the set threshold (default 0.3). If the stopping point (left boundary) of the sliding windows is more
710 than 300 bp away from the left end of the consensus sequence, TEtrimmer regards the left end sequence
711 extension as complete. Otherwise, a new round of extension is initiated.

712 When the left end sequence extension is finished, TEtrimmer conducts right end sequence extension of
713 the MSA. Similar to the steps described above, right end sequence extension also only aligns extended
714 sequences from the current round and generates the respective consensus sequence. In contrast to the
715 left end, to judge if the overall right end extension is sufficient, a right end sliding window is created
716 only for this consensus sequence. The sliding window stops when the proportion of the ambiguous letter
717 "N" (as defined above) within the sliding window is less than the set threshold (default 0.3). If the
718 stopping point (right boundary) of the sliding window is more than 300 bp away from the right end of
719 the consensus sequence, the right end extension is complete.

720 Note that this way extension sizes for the two ends of each sequence in the MSA may differ. After
721 sequence extension at both ends, TEtrimmer calculates the new (extended) genome coordinates
722 (locations) for each sequence. Finaly, extended sequences are extracted from the corresponding genome.
723 All sequences are aligned and gappy columns are removed from the MSA by the function
724 "`remove_gap_columns`" as described above (see section "MSA cleaning").

725 **TE boundary definition**

726 Sequence extension helps to complete TE consensus sequences, but putative excessive extension can
727 also add other, non-TE sequences (false-positives) at both ends of the MSA. TEtrimmer uses two
728 strategies to identify TE boundaries and to remove such inappropriate regions. Based on the final MSA
729 after the extension step, first a consensus sequence is generated for each MSA, and "N" is used to
730 represent any ambiguous column where the proportion of the predominant nucleotide is below 0.8 or
731 the total count of nucleotides is less than five. Then, a self-BLASTN search (with relatively relaxed e-
732 value of 0.05) of the consensus sequence is conducted to identify potential terminal repeats of TEs,
733 including LTRs and TIRs. Once a terminal repeat is found, the TE boundary is set at the distal position
734 of the terminal repeat, and the MSA is cropped accordingly. On the other hand, if no terminal repeat is
735 identified, TEtrimmer selects sliding windows (default size 150 bp) at both ends of the consensus
736 sequence that stepwise move towards the opposite end of the sequence nucleotide position by nucleotide
737 position until the proportion of the ambiguous letter "N" is below 0.2 within the sliding windows. Then,
738 only the MSA columns between the stopping points of the two sliding windows (including the sliding
739 windows) are kept. This procedure removes the majority of false-positive flanking sequences.
740 Furthermore, the functions "`crop_end_by_divergence`" and "`crop_end_by_gap`" (see
741 section "MSA cleaning above) remove highly divergent regions from the MSA. Based on the defined
742 TE boundaries, TEtrimmer generates the corresponding consensus sequence for further analysis.

743 **ORF and PFAM domain prediction**

744 For autonomous TEs, the presence of characteristic protein domains can be used to assist TE
745 classification (Goubert et al., 2022) and to determine TE orientation. TEtrimmer adopts the PFAM
746 database (Mistry et al., 2021) to annotate protein domains within TEs. It either uses the "`--pfam_dir`"
747 option to query a local PFAM library, or it can download the current PFAM database automatically in

748 the absence of a local PFAM instance. Thereafter, TEtrimmer uses the EMBOSS function "`getorf`"
749 (Rice et al., 2000) to predict ORFs and "`pfam_scan.pl`" (Mistry et al., 2021) to search for TE protein
750 domains against the PFAM database. Typically, all predicted PFAM domains share the same orientation.
751 TEtrimmer generates reverse complements of the TE consensus sequence and the MSA if all PFAM
752 domains are found on the antisense strand of the consensus sequence. If TE-related PFAM domains are
753 found on both strands, TEtrimmer sums the length of PFAM domains found on both strands to decide
754 on the orientation of the TE consensus sequence via a majority decision.

755 **TE classification**

756 TEtrimmer includes two TE classification methods. First, it adopts the RepeatModeler2 module
757 "`RepeatClassifier`" (Flynn et al., 2020) to categorize TEs. By default, TEtrimmer re-assigns
758 processed TEs if the sequence extension size is larger than 4000 bp as these may represent mis-
759 categorized and/or fragmented TEs. Further, two options are associated with the first classification
760 method, named "`--classify_all`" and "`--classify_unknown`". TEtrimmer runs
761 RepeatClassifier for each output if the "`--classify_all`" option is used and only re-classifies
762 "Unknown" TEs via the "`--classify_unknown`" option. If RepeatClassifier assigns a new
763 category, TEtrimmer will replace the old assignment with it; alternatively, the original classification is
764 kept. The second method is performed when the initial analysis of all the input TE sequences is
765 completed. TEtrimmer uses the final, classified output sequences as a database to re-assign unknown
766 sequences via RepeatMasker.

767 **Output evaluation**

768 Five different evaluation levels including "Perfect", "Good", "Reco_check" (i.e., recommended check),
769 "Need_check", and "Low_copy" are assigned to each TEtrimmer output. TEtrimmer evaluates TE
770 consensus sequences based on the presence of terminal repeats, classification status, the number of
771 sequences in the MSA, the number of full-length BLASTN hits, and PFAM protein domain predictions
772 (Table 1). A full-length BLASTN hit is defined as a sequence with more than 90% coverage and more
773 than 85% identity with the respective query sequence (i.e., the TEtrimmer-processed consensus
774 sequence). When the number of sequences within the MSA is less than ten, TEtrimmer examines if the
775 input sequence contained terminal repeats and if the number of full-length BLASTN hits is greater than
776 or equal to two. If so, this input sequence is regarded as a low-copy element, otherwise it is excluded
777 from further analysis.

778 **TE consensus library de-duplication**

779 All consensus sequences processed by TEtrimmer are stored into the file "TEtrimmer_consensus.fasta",
780 which potentially contains duplicated sequences. To reduce any putative redundance, TEtrimmer
781 performs two rounds of de-duplication by CD-HIT-EST (Fu et al., 2012). For the first round, consensus
782 sequences are grouped into one cluster if they share more than 90% sequence identity and the alignment
783 coverage of the shorter sequence is more than 90%. Afterwards, TEtrimmer selects all clusters that
784 contain at least one "Perfect" or "Good" (Table 1) consensus sequence. For each selected cluster, if
785 "Perfect" candidates exist, the longest "Perfect" consensus sequence is selected as the representative. If
786 only "Good" candidates are present, the longest "Good" consensus sequence is selected. The second
787 de-duplication round extracts all sequences from the remaining clusters and uses these sequences again
788 as input for CD-HIT-EST. Sequences are grouped into one cluster if they have more than 85% identity
789 and the alignment coverage for all sequences is greater than 80%. Only the longest sequence in each
790 new cluster is selected for the de-duplicated TE consensus library. The de-duplicated TE consensus
791 sequences are saved into the text file "TEtrimmer_consensus_merged.fasta".

792 **Generation of report plots**

793 Seven plots are generated for each output to help evaluating the TEtrimmer output and to compare the
794 differences between the TEtrimmer-processed and input consensus sequences. The report plots file
795 include (1) an MSA ends plot, (2) a whole MSA plot, (3 and 4) TE-Aid (Goubert et al., 2022) plots for
796 both input and output sequences, (5 and 6) a PFAM plot for both input and output sequences, and (7) a
797 dot plot comparing input and output sequences (Fig. 7). For the MSA ends plot, TEtrimmer extracts 50
798 columns before and after the identified TE boundaries (see section "TE boundary definition" above)
799 from the MSA and artifactually joins them by ten hyphen symbols, signifying the TE (Fig. 4A).
800 Nucleotides are represented by different colors, and the nucleotide background is colored if the
801 nucleotide proportion in the column is less than 40%. The whole MSA plot module is derived from the
802 software CIAlign (Tumescheit et al., 2022) and the nucleotides are represented by different colored bars
803 (Fig. 4B). In contrast to the default CIAlign plot, TEtrimmer adds two arrows to indicate the TE
804 boundaries (TE start and TE end) in the whole MSA plot. TEtrimmer also includes the TE-Aid package
805 to help plotting. All TEtrimmer output and input TE consensus sequences are used for TE-Aid analysis
806 (Fig. 4C and D). Furthermore, the PFAM plots are generated based on ORF and PFAM domain
807 prediction results, which are denoted as blue and red arrows, respectively (Fig. 4E). The last dot plot is
808 generated by the EMBOSS tool "`dotmatcher`" (Rice et al., 2000). The position of the consensus
809 output sequence after analysis by TEtrimmer is plotted on the x-axis and the position of the
810 corresponding input TE consensus sequence is plotted on the y-axis of the dot plot (Fig. 4F). All seven
811 report plots are merged into one file to be used for convenient output evaluation.

812 **Used genomes and reference TE consensus libraries for genome-wide benchmarking**

813 The genomes of six eukaryotic organisms were used to test the TE annotation performance of
814 TEtrimmer at the level of TE consensus libraries and a genome-wide scale. These species comprised *B.*
815 *hordei* (barley powdery mildew fungus), *D. melanogaster* (fruit fly), *D. rerio* (zebrafish), *O. sativa*
816 (rice), *Z. mays* (maize), and *H. sapiens* (human) (Table 1). The reference TE consensus library of *B.*
817 *hordei* is based on in-house manually curated TEs (Qian et al., 2023). In case of *D. melanogaster*, *D.*
818 *rerio*, *O. sativa*, and *Z. mays*, Repbase 28.10 (Weidong. Bao et al., 2015) was used for generating the
819 respective reference TE consensus libraries. *H. sapiens* reference TE consensus library was extracted
820 from Dfam (Hubley et al., 2016).

821 **Benchmarking of MSA cleaning performance**

822 Due to the presence of fragmented TEs and/or excessive end extension, the extended MSAs typically
823 contain regions of low sequence conservation, which can be efficiently cleaned by the TEtrimmer
824 cleaning module. The MSA cleaning capability of TEtrimmer was evaluated quantitatively by a
825 customized confusion matrix (Provost et al., 1998). Three MSAs were used for each evaluation,
826 including the original MSA before cleaning, the manually cleaned MSA, and the MSA cleaned by
827 TEtrimmer. The manually cleaned MSA served as the reference in the benchmarking process.

828 Both manual cleaning and TEtrimmer-based cleaning did not delete the nucleotides in lowly conserved
829 (poorly aligned) regions but converted them to gaps (-). Consequently, all three MSAs had the same
830 number of columns and sequences. The order of the sequences within the MSA was also identical.
831 Thereafter, each position in the three MSAs was analyzed and corresponding true positive (TP) sites,
832 false negative (FN) sites, false positive (FP) sites, and true negative (TN) sites values were calculated
833 and summed into TP, FN, FP, and TN values (Table 3). For example, if a site in the original MSA was
834 occupied by any type of nucleotide, this site was counted as "true positive" when both manual and
835 TEtrimmer-based cleaning converted this site to a gap (-). At the end, the TEtrimmer cleaning sensitivity

836 (TP / (TP + FN)), precision (TP / (TP + FP)), and F1 score (2 * (sensitivity * precision) / (sensitivity +
837 precision)) were calculated and used to evaluate the TEtrimmer cleaning performance.

838 **Table 3. Confusion matrix calculation for the MSA cleaning performance.**

	True positive	False negative	False positive	True negative
Original MSA	Nucleotide	Nucleotide	Nucleotide	Nucleotide
Manually cleaned MSA	Gap	Gap	Nucleotide	Nucleotide
TEtrimmer-cleaned MSA	Gap	Nucleotide	Gap	Nucleotide

839

840 **Benchmarking of genome-wide TE discovery and annotation**

841 We benchmarked TEtrimmer, RepeatModeler2, and EDTA2 on the basis of the corresponding reference
842 TE libraries (Table 1) by two methods. The first method directly compared the consensus sequences to
843 assess the integrity of TE sequences, which aimed to determine how many TE sequences from the
844 reference library can be recovered by the respective de novo-generated library. The other method
845 compared genome-wide TE annotation by the different tools to evaluate if TEs in the genome were
846 correctly masked. Six organisms were used for the benchmarking test and the reference TE consensus
847 libraries and genomes were downloaded and prepared (Table 1). RepeatModeler2 (Flynn et al., 2020)
848 with parameters "`-threads 50 -LTRStruct`" and EDTA2
849 (https://github.com/oushujun/EDTA/tree/EDTA2) with parameters "`--threads 50 --`
850 `sensitive 1`" were run to generate de novo TE libraries for all selected organisms. Thereafter, de
851 novo-generated TE consensus libraries were used as the input for TEtrimmer analysis with parameters
852 "`--num_threads 50 --classify_all`". Finally, all TE consensus libraries generated before
853 and after TEtrimmer analysis were used individually to annotate the corresponding genome by
854 RepeatMasker v4.1.1 with parameters "`-pa 50 -s -a -inv -gff`".

855 First, we performed benchmarking by direct comparison of the TE consensus libraries. We used the
856 benchmarking method developed by Flynn and co-workers (Flynn et al., 2020) for this purpose. All TE
857 consensus libraries generated by EDTA2, RepeatModeler2, and TEtrimmer were compared with the
858 corresponding reference TE consensus library (Table 1). For EDTA2, we created an additional library
859 for the benchmarking by combing the EDTA2 consensus library with all TE sequences identified by
860 the EDTA2 structure-based module. This new EDTA2 library was then de-duplicated using CD-EST-
861 EST with the parameters "`-c 0.9 -aL 0.9 -aS 0.9`", resulting in the new library named "EDTA2
862 struct-lib". Due to the default separation of flanking LTRs from internal sequences (LTR-INT) in the
863 case of LTR retrotransposon by tools like EDTA2 and RepeatModeler2, as well as in reference TE
864 consensus libraries, a custom Python script was developed to separate flanking LTRs from the internal
865 sequences (LTR-INT) for all TEtrimmer consensus libraries. The respective script was named
866 "SeparateLTR.py", which can be downloaded from the TEtrimmer GitHub repository. To avoid the
867 benchmarking bias caused by flanking LTRs, we did not include the flanking LTRs for benchmarking.
868 Four evaluation levels were assigned to each TE sequence in the reference library, including "Perfect",
869 "Good", "Present", and "Not found". "Perfect" means that the reference TE consensus sequence has
870 one match in the de novo-generated library with > 95% similarity and coverage. "Good" indicates that
871 the reference TE consensus sequence has multiple overlapping matches in the de novo-generated
872 library, each with > 95% similarity and total coverage. "Present" is similar to "Good", but the required
873 minimum similarity and total coverage is decreased to 80%.

874 Next, we performed the benchmarking at the level of genome-wide TE annotation. We developed a
875 custom python script named "TE_sankey_plotter.py" (available from TEtrimmer GitHub repository),

which employs a confusion matrix to evaluate the genome-wide TE annotation and to illustrate the outcome comparatively via Sankey plots. The details about how to apply a confusion matrix on the genome-wide TE annotation was described before (Ou et al., 2019). Briefly, the RepeatMasker ".out" annotation files derived from the TEtrimmer, RepeatModeler2, and EDTA2 consensus libraries, respectively, were compared with the RepeatMasker ".out" file based on the reference TE consensus library. First, all ".out" files were sequentially converted into BED files with four columns (chromosome name, start position, end position, and TE type). Then, the overlapping TE regions were resolved by BEDtools v2.31.1 (Quinlan & Hall, 2010) with parameters "`-d 0 -c 4,4,2,3 -o collapse, distinct, collapse, collapse`". Finally, six metrics scores were calculated, i.e., sensitivity, specificity, accuracy, precision, F1 score, and false discovery rate, to evaluate the genome-wide TE annotation.

Computational resource requirements and installation of TEtrimmer

TEtrimmer is written in the language Python3 and can be conveniently installed via a Conda package or a Docker image at https://github.com/qjiangzhao/TEtrimmer. TEtrimmer can be run on the operational systems Linux, macOS, and a Windows subsystem for Linux (WSL). There are no specific hardware requirements for running TEtrimmer. TEtrimmer is multi-threaded tool and supports high-performance computing (HPC).

Abbreviations

bp	base pair
DBSCAN	Density-Based Spatial Clustering of Applications with Noise
EDTA2	Extensive De novo Transposon Annotation 2
GUI	Graphical user interface
LINE	Long interspersed nuclear element
LTR	Long terminal repeat
LTR-INT	Long terminal repeat internal sequence
MSA	Multiple sequence alignment
ORF	Open reading frame
PCA	Principal component analysis
PFAM	Protein families database
SINE	Short interspersed nuclear element
TE	Transposable element
TIR	Terminal inverted repeat

Declarations

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Author contributions

RP, SK, and JQ conceived this study.
JQ and HX developed the TEtrimmer software.
SK performed proofreading of the TEtrimmer code.
HX and JQ drafted the TEtrimmer manual.
LF supported the development of the TEtrimmer MSA cleaning procedure and the MSA clustering algorithm.
SO provided EDTA2 results and performed intensive de-bugging of TEtrimmer.
JS conducted RepeatMasker analysis.
JQ and SK wrote the first draft of the manuscript.
JQ drafted the Figures.
MW supervised work performed by HX and provided funding.
RP and SK supervised work performed by JQ.
RP provided funding.
All authors reviewed the manuscript and contributed to the editing.

Competing interests

The authors declare they have no competing interests.

Ethics approval

Not applicable.

Data availability

TEtrimmer is an open-source software under license GPLv3. We used large language models ChatGPT and GPT4 to facilitate the development of TEtrimmer. The code can be downloaded from GitHub at https://github.com/qjiangzhao/TEtrimmer. A detailed TEtrimmer manual is supplied. All benchmarking results and scripts can be found at GitHub at https://github.com/qjiangzhao/TEtrimmerPaperFile. All third-party software included in the TEtrimmer package are published under open-source licenses, with the exception of TE-Aid, for which we have approval by the author Clément Goubert (University of Arizona, USA).

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Supplementary files

Supplementary Figure 1. Multiple sequence dot plot based on all sequences in a TE cluster.

Supplementary Figure 2. The TEtrimmer GUI application can help to inspect and improve any TE consensus library.

Supplementary Table 1. Relative tree branch distance matrix of the un-rooted maximum-likelihood phylogenetic tree calculated on the basis of the divergent columns in the MSA.

Supplementary Table 2. An exemplary report table generated by TEtrimmer.

Supplementary Table 3. Number of TE sequences in each benchmarking category in comparison to the respective reference TE consensus library (raw data for Fig. 6A).

Supplementary Table 4. Types of TEs in the "Perfect" benchmarking class (raw data for Fig. 6B).

Supplementary Table 5. Confusion matrix scores related to the genome-wide benchmarking.

Supplementary File 1. Example of interactive report plots generated by the GUI application.