METHOD

TransposonUltimate: software for transposon classification, annotation and detection

Kevin Riehl¹, Cristian Riccio^{1,2}, Eric A. Miska^{1,2,3} and Martin Hemberg^{2,4*}

*Correspondence:

mhemberg@bwh.harvard.edu

⁴Evergrande Center for

Immunologic Diseases, Harvard Medical School and Brigham and Women's Hospital, 75 Francis Street, MA 02215 Boston, MA, USA

Full list of author information is available at the end of the article

†Equal contributor

Abstract

Motivation: Most genomes harbor a large number of transposons, and they play an important role in evolution and gene regulation. They are also of interest to clinicians as they are involved in several diseases, including cancer and neurodegeneration. Although several methods for transposon identification are available, they are often highly specialised towards specific tasks or classes of transposons, and they lack common standards such as a unified taxonomy scheme and output file format. Moreover, many methods are difficult to install, poorly documented, and difficult to reproduce.

Results: We present TransposonUltimate, a powerful bundle of three modules for transposon classification, annotation, and detection of transposition events. TransposonUltimate comes as a Conda package under the GPL-3.0 licence, is well documented and it is easy to install. We benchmark the classification module on the large *TransposonDB* covering over 891,051 sequences to demonstrate that it outperforms the currently best existing solutions. The annotation and detection modules combine sixteen existing softwares, and we illustrate its use by annotating *Caenorhabditis elegans*, *Rhizophagus irregularis* and *Oryza sativa subs. japonica* genomes. Finally, we use the detection module to discover 29,554 transposition events in the genomes of twenty wild type strains of *Caenorhabditis elegans*.

Availability: Running software and source code available on <https://github.com/DerKevinRiehl/TransposonClassifierRFSB>. Databases, assemblies, annotations and further findings can be downloaded from <https://cellgeni.cog.sanger.ac.uk/browser.html?shared=transposonultimate>.

Keywords: Transposable elements; Transposon classification; Transposon annotation; Transposon detection

1 Introduction

Transposons are evolutionary ancient mobile genetic elements that can move via copy&paste and cut&paste transposition mechanisms. They can be classified within a taxonomic scheme (Fig. 1A), and each class is associated with a set of characteristics, e.g. proteins relevant for transposition and structural features (Fig. 1B). During transposition, transposable elements (TEs) can leave structural patterns both at the insertion and the deletion site [1, 2, 3]. Autonomous transposons encode the tools necessary for transposition events, e.g. genes producing transposase, integrase and other enzymes [3], while non-autonomous transposons depend on proteins encoded elsewhere [4]. As the insertion of a transposon can be detrimental, many species have developed repression mechanisms, e.g. TE promoter methylation [5] and piRNAs [6]. Even though transposition events occur rarely [7], in many

organisms large sections of DNA consist of either transposons or their transposition-incompetent descendants that have accumulated mutations over time [4]. It is estimated that transposons make up a large share of the genome in many species; 45% in humans, 20% in fruit flies, 40% in mice, 77% in frogs and 85% in maize [8].

Studying TEs is highly relevant for understanding evolutionary processes [9], developmental biology, gene regulation, and many diseases called transposonopathies such as subtypes of haemophilia, immunodeficiency, cancer and Alzheimer's disease [10, 11, 12]. Also, TEs are popular for genetic engineering purposes as they allow for direct insertion of their genetic cargo into a target genome [13, 14, 15]. However, the repetitive nature of transposons and their descendants is a challenge for their analysis and discovery, in particular when using short-read sequencing technologies [7]. Long-read technologies facilitate studies of transposons and their functional consequences, but they also require novel computational tools. Although various approaches for identifying transposons have been proposed recently [16], current tools are error prone, not robust, mostly rely on prior knowledge of transposon sequences, and are often limited to a family of transposons or a group of species [17].

Here, we present a bundle of tools addressing three different tasks related to transposon identification: classification, annotation and detection. The goal of classification is to determine which taxonomic class a given transposon sequence belongs to. The annotation task consists of scanning a genome sequence to mark all transposons. Finally, the detection task involves the comparison of two genomes to identify structural variants arising from the insertion of TEs.

Existing transposon classifiers are difficult to compare directly since they vary in their approach, which features and taxonomies they use, how they evaluate predictions, and which databases are used for training. Applications of SVMs [18], hidden Markov models [19], random forests [20], Gaussian naive Bayes [21], decision trees [22], stacking [23, 24], boosting [25, 26], neural networks [27, 28, 29], evolutionary algorithms [30, 21] and genetic algorithms [31, 32, 33, 34] can be found in the literature. Most methods use sequence features, such as the k-mer frequency, the occurrence of structural [35] and protein features [18] for classification. Besides, another approach is to classify TEs using the similarity to known transposons based on a sequence library [36].

The annotation of transposons in nucleotide sequences is challenging due to the presence of transposition-incompetent TEs that have been mutated, truncated, degraded, fragmented and dismembered due to nesting [37]. Annotation is further complicated by a lack of standards [38] and disagreement on definition, taxonomy and terminology [39, 40]. Since transposons do not adhere to a universal structure [41], many researchers have employed class-specific approaches [42]. Moreover, most of the software employed for transposon annotation was originally designed for gene annotation, neglecting the peculiarities of transposons [39]. Existing transposon annotation methods (Table 1) can be assigned to one or more approaches [43, 2, 1, 41]. The *de novo* approach finds transposons by identifying repetitive sequences. It is effective in discovering previously unknown transposons with high prevalence [41], but it is computationally costly [41, 39], unable to find degraded transposons [41], and risks misidentifying repetitive DNA or high copy

number genes as transposons [44, 45]. The structure-based approach (also called motif-based [42] or signature-based approach [2]) is based on knowledge of the structure of transposons and annotates by finding combinations of characteristic patterns [38, 46]. This approach enables the discovery of transposition-incompetent transposons thanks to their unique structural properties [41]. However, these approaches are often characterised by high false discovery rates [37, 44] and they miss transposons with weak signatures [37]. The similarity-based approach (also called library-based approach [2]) employs a library of known transposons together with BLAST(-like) tools. The high accuracy [41] and short runtimes [44, 47] of this approach come at the cost of its inability to find unrelated transposons [41, 47] and the dependency on quality and exhaustiveness of the library [38, 44, 48]. Moreover, the current version of the most widely used database RepBase [49] is behind a paywall and the related tools RepeatMasker and RepeatModeler are not transparent with regards to how transposons were curated and consensus sequences were generated [39].

Previous efforts to detect transposition events by comparing two genomes have been based on the analysis of the depth of coverage, discordant and split read pairs [50, 51]. However, both the task of detecting structural variants (SVs) and annotating TEs are very challenging when using short reads [7]. Recently, long-reads technologies have become more widely available, but to the best of our knowledge the only existing method that can take advantage of them for TE detection is LoRTE [52]. Although results indicate that LoRTE performs well even on low coverage reads, it is limited to PacBio data and insertion and deletion SVs only.

Here, we present TransposonUltimate, a set of tools for the identification of transposons, consisting of three modules for accurate classification, annotation in nucleotide sequences and detection of transposition events (Fig. 2). Our new classifier is benchmarked against existing softwares, and we use the annotation module to analyse the genomes of three different species. Finally, the detection module is employed to identify transposition events in 20 high quality genomes from *Caenorhabditis elegans* wild isolates that were assembled using a combination of long- and short-read technologies.

2 Materials and methods

2.1 Transposon classification module, RFSB

Given a nucleotide sequence that is considered to be a transposon, the goal is to determine the class of a transposon according to a given taxonomy. This task is a hierarchical classification problem, meaning the classifier needs to identify multiple classes that stand in a relationship described by a taxonomic hierarchy. The design of the classification module includes several aspects; choosing a transposon database, feature selection, model structure, training strategy, model implementation, evaluation and benchmarking.

The classifiers considered here are supervised learning algorithms, and consequently their performance is limited by the data used for training. Previous studies used small transposon sequence databases, each with different taxonomic schemes, which does not allow for a direct comparison. Therefore, we created *TransposonDB* (Fig. 3, File F1), a large collection of transposon sequences that consists of ten

databases: ConTEdb [53] (<http://genedenovoweb.ticp.net:81/conTEdb/index.php>), DPTEdb [54] (http://genedenovoweb.ticp.net:81/DPTedB/browse.php?species=cpa&name=Carica_papaya_L.), mipsREdat-PGSB [55] (<https://pgsb.helmholtz-muenchen.de/plant/recat/index.jsp>), MnTEdb [56] (<http://genedenovoweb.ticp.net:81/MnTEdb1/>), PMITEdb [57] (http://pmite.hzau.edu.cn/download_mite/), RepBase [58] (<https://www.girinst.org/repbase/>)^[1], RiTE [59] (<https://www.genome.arizona.edu/cgi-bin/rite/index.cgi>), Soytedb [60] (<https://www.soybase.org/soytedb/#bulk>), SPTEdb [61] (http://genedenovoweb.ticp.net:81/SPTEdb/browse.php?species=ptr&name=Populus_trichocarpa) and TrepDB [62] (<http://botserv2.uzh.ch/kelldata/trep-db/downloadFiles.html>). To create the database, the taxonomies were unified, duplicates were dropped and several filter rules were applied (Table S1). Filtering included the removal of sequences with no label, the exclusion of fragments, contigs, satellites and RNA sequences. Moreover, only sequences with a length greater than 100bp and those including at least once each of the letters 'A', 'C', 'G' and 'T' were kept. To the best of our knowledge, this is the largest database of transposon sequences available. Since TransposonDB covers all relevant Eukaryotic kingdoms, it allows for the training and evaluation of a robust, cross-species hierarchical classification model (Table S2 + S3). Moreover, the database is balanced and covers sufficient examples for all taxonomic nodes (Table S4).

We selected the combination of relative k-mer frequencies and binary protein features for our classifier. Relative k-mer frequencies represent the number of occurrences of a k-mer within a sequence divided by the number of times it would appear if the sequence consisted of this k-mer only. Protein features are binary, indicating the presence of a certain protein domain in the sequence. The feature vector consists of k-mer frequencies ($k = 2, 3, 4$) and 169 selected domains from NCBI CDD [63] covering class-specific transposons (Table S5). RPSTBLASTN (v2.10.1) was used to annotate the conserved domain models at an e-value of 5.0 as it performed best in terms of classification performance (Fig. S1 A-B). In addition, two model structures were explored. The binary structure employs binary classifiers for each node (= transposon class) of the taxonomy. After inference of the binary classifiers, the taxonomic class can be determined by choosing the most probable node at each stage. The multilabel structure employs a multilabel classifier for each parent node of the taxonomy with $n+1$ classes representing the taxonomic child classes and -1 (return scenario). After inference, the taxonomic class can be determined by choosing the most probable child node at each stage or to return to a higher level and then choose the second most probable child node at that stage. Moreover, we explored two training strategies. The comprehensive training strategy trains each classification node with the whole training set, while the selective training strategy trains each classification child node with a training set that was activated by the parent node. All training strategies, model structures and feature generation were implemented in Python (v3.6.9). Models implementing random forests, AdaBoost, logistic regression, SVM and Naive Bayes from the machine learning package scikit-learn (v0.23) [64] were explored. Random forest consistently

^[1]We use version 23.08 that was the last publicly available version before the paywall was introduced.

yields the highest classification performance (Fig. S2). Based on these results, we propose a random forest classifier with a selective training strategy on a binary model structure, *RFSB*.

Previous transposon classification studies use different performance measures, taxonomies, training and testing sets, making it hard to compare them. To evaluate the performance, we consider three perspectives. The first perspective is based on hierarchical precision and recall, meaning it considers the whole taxonomy, as proposed in [65]. The second perspective evaluates for different taxonomic levels and the third perspective captures the classification performance of single classes. We benchmark RFSB againsts TERL [29], TopDown [24], NLLCPN [27], HC_LGA [33] and HC_GA [31], as their published code allowed for reproduction. To ensure a fair comparison, source codes were partially modified to allow the training and evaluation of these models on the taxonomy used in our work and TransposonDB.

2.2 Transposon annotation module, *reasonaTE*

Given an assembled genome, the goal of the annotation module is to find all transposon occurrences and their locations. Our *reasonaTE* pipeline produces rich annotations, including transposon mask regions (union of all annotated base pairs) as well as transposon annotations, classification, structural and protein features. This is achieved by combining the advantages of thirteen published transposon annotation tools covering different annotation approaches and transposon classes: RepeatMasker (<http://www.repeatmasker.org/>), RepeatModeler (<http://www.repeatmasker.org/RepeatModeler/>), LTRharvest [66] (<https://www.zbh.uni-hamburg.de/forschung/gi/software/ltrharvest.html>) and TIRvish [67] (http://genometools.org/tools/gt_tirvish.html) are available as Conda packages. Moreover, we created Conda packages for SINE-Finder [68] (http://www.plantcell.org/content/suppl/2011/08/29/tpc.111.088682.DC1/Supplemental_Data_Set_1-sine_finder.txt), SINE-Scan [69] (https://github.com/maohlzj/SINE_Scan), HelitronScanner [42] (<https://sourceforge.net/projects/helitronscanner/files/>), MUSTv2 [70] (<http://www.healthinformatics.org/supp/resources.php>), MiteFinderII [71] (<https://github.com/jhu99/miteFinder>) and MITE-Tracker [72] (<https://github.com/INTABiotechMJ/MITE-Tracker>) to make them accessible and to facilitate their installation. Also, we include the output files of LTRpred [73] (<https://hajkd.github.io/LTRpred/articles/Introduction.html>) into the pipeline, as this tool provides high quality annotations, but is available as a Docker image only. As the tools have different output formats, we developed a parser module to convert all outputs to GFF3 format.

After running the annotation tools, additional copies of the identified transposons are searched using the clustering tool CD-HIT (v4.8.1) [74, 75] and BLASTN (v2.10.1). For the annotation of transposon-characteristic proteins, we have created a Conda packaged version of TransposonPSI (<http://transposonpsi.sourceforge.net/>), and we also use the protein domains from NCBI CDD for this task. Using TransposonDB, NCBI CDD and RPSTBLASTN, we selected the 1,000 most frequently occurring protein domains that are characteristic to transposons (File F2). As an application, here we annotate the genome *MSU7* of *Oryza sativa subspecies japonica* (<http://rice.plantbiology.msu.edu/index.shtml>),

the genome *DAOM197198* of *Rhizophagus irregularis* (<https://www.ncbi.nlm.nih.gov/bioproject/?term=PRJDB4945>) [76], three reference genomes *VC2010* (<https://www.ncbi.nlm.nih.gov/bioproject/?term=PRJEB28388>), *N2* (<https://www.ncbi.nlm.nih.gov/bioproject/?term=PRJNA13758>), *CB4856* (<https://www.ncbi.nlm.nih.gov/bioproject/?term=PRJNA275000>) and twenty novel wild type strains [77] of *Caenorhabditis elegans* (Table S6).

2.3 Transposition event detection module, deTEct

Given an assembled reference genome and sequenced probe genome reads, the goal is to identify transposition events that are manifested as structural variants. This requires both a list of SVs and annotation of TEs as inputs. We employ the structural variant caller Sniffles on ngmlr [78] alignments and PBSV (<https://github.com/PacificBiosciences/pbsv>) structural variant caller on pbmm2 alignments (<https://github.com/PacificBiosciences/pbmm2>). Moreover, the TE annotations are generated using the proposed reasonaTE pipeline mentioned before.

SVs are filtered twice. First, variants shorter than 50 bp or longer than 1% of the genome length were excluded. Second, duplicate structural variants of the same type are merged. Consecutively, the remaining variants and TE annotations are matched and finally reported if their length corresponds to each other. Transposon annotations were matched to structural variants if they intersected for at least 10% and their length was similar by a threshold of 50%. We chose to do so, as structural variant callers and transposon annotators have an uncertainty regarding exact locations. We therefore consider a similar length more important than a high overlap. The proposed deTEct pipeline is applicable to long-read sequencing technologies, and it has been tested with both PacBio and OxfordNanopore data.

3 Results

3.1 RFSB outperforms other transposon classifiers

We benchmarked our RFSB method against other transposon classifiers, and the results show that it has the highest sensitivity and specificity (Fig. 4 A, Table S7). TE Learner [20] has the lowest reported performance, while the other methods have similar F1 scores. However, this comparison is based on reported numbers from different studies with different evaluation schemes, taxonomies and datasets for training and testing. For a more fair comparison some of the tools were applied to the subset of TransposonDB which includes RepBase and PGSB (Fig. 4 B). The comparison of the results reveals large discrepancies. Surprisingly, TERL and TopDown have a performance which is worse than random guessing, and closer inspection of the outputs from NLLCPN reveals that it has learned a constant distribution rather than a relationship between sequences and classes.

A detailed analysis of the classification performance of RFSB across different taxonomic levels and classes reveals a small decrease in performance when considering deeper taxonomic levels (Fig. 4 C). Underrepresented classes, e.g. Helitrons and MITEs, perform worse, and the results are consistent for both F1 and MCC scores. Moreover, for some classes the performance of RFSB on the large, cross-species TransposonDB is better than for the more homogeneous subset of RepBase and PGSB, which suggests that it is robust, generalisable, and applicable to different

species. An inspection of the most informative features (File **F3**) shows that longer k-mer features contribute the most to the classification performance, while protein domains have a smaller share amongst the most contributing features (Fig. 4 D). This motivated the exploration of longer k-mer features, but we did not find any significant increase of the performance when using 5-mers (Fig. 4 E).

3.2 The ensemble strategy reasonaTE finds more transposons and reduces bias

Next, we evaluated the ability of our reasonaTE pipeline to identify TEs in the genomes of three different species (Fig. 5 A-B, File **F4**). The TE content of almost 21% for *Caenorhabditis elegans* is higher than previously reported values of 12% [8], 17% [79], and 12-16% [80]. However, as these studies used methods that were biased towards finding specific classes of transposons, it is to be expected that our ensemble strategy finds more TEs. By contrast, the prediction of 33% for *Oryza sativa subs. japonica* is very close to the mean of other reports [81, 82, 83, 84, 85, 86, 87, 88, 89]. The content of 23% in *Rhizophagus irregularis* is close to a previous estimate of 27% [90]. The low variation of transposon content across different strains becomes obvious for the cluster of *Caenorhabditis elegans*. Interestingly, the relative transposon class frequency reveals clear differences across species (Fig. 5 C-D). Similarly, the length distributions (Fig. 5 E-G) exhibit substantial differences between transposons of the same class found in different species. Helitrons in particular vary in length as was observed before [91].

In concordance with [92] and [93], the share of Helitrons amounts to almost 2% of the *Caenorhabditis elegans* genome. Moreover, the majority of the transposons are TIR DNA transposons, as reported by [94, 95, 79]. Contrary to previous studies [80, 96, 97], we mainly find hAT, CMC and Novosib transposons to be present in the *Caenorhabditis elegans* genome rather than Tc1-Mariner transposons. Our findings for the rice genome are consistent with previous findings. The high frequency of Gypsy (class 1/1/2) compared to other LTR (class 1/1) and non-LTR (class 1/2) was reported in *Oryza sativa subs. japonica* [87]. Moreover, the small share of MITEs, up to 2%, is similar to the previously reported share of 4% [89]. A previous study [44] found that class 1 transposons have a larger share (25%) than class 2 transposons (20%) and the frequencies for the subclass level (LTR 23.5% and non-LTR 2%, TIR 17.5% and Helitrons 3.6%) match our findings. Inspection of the annotation density across the chromosomes revealed a characteristic concentration at the arms for *Caenorhabditis elegans* (Figure S3), consistent with the higher densities observed for other variants [79, 98, 99, 100, 101, 80].

The comparison of different annotation tools reveals that reasonaTE provides more unbiased results (Figure S4) as none of the other methods find more than 31.8% of the TEs reported by reasonaTE. In addition, the analysis shows that around 40% of the repetitive elements found by RepeatMasker and RepeatModeler were confirmed as transposons using our approach. Moreover, the transposon characteristic protein annotations by TransposonPSI and the 1,000 most frequently occurring proteins from NCBI CDD intersect significantly with reasonaTE's transposon annotations. The analysis also reveals large overlap between some tools, e.g. MUSTv2 & MITE-Tracker, LTRpred & LTRharvest, and SINE-Finder with all other tools.

Closer inspection of the class composition of the TEs found for *Caenorhabditis elegans* confirms the advantages of the ensemble technique of reasonaTE (Figure S5). None of the tools is able to find the same share of TEs on its own as the ensemble. Moreover, we find that tools that were designed to identify a specific transposon class annotate TEs from different classes as well.

3.3 29,554 transposition event candidates were observed analyzing 20 wild type strains of *Caenorhabditis elegans* using deTEct

Finally, we applied the deTEct pipeline to 20 whole genome assemblies of wild type strains of the nematode *Caenorhabditis elegans*. Each strain was compared to the two reference genomes *VC2010* and *CB4856* (Fig. 6 A, Table S8, File F5). As expected, the newly sequenced genomes of these two strains have almost no transposition events when compared to their reference. Closer inspection of the transposon and transposition event densities reveals that the putative transposition events are primarily located at the ends of the chromosomes (Fig. 6 B) as reported by [79]. From the initial list of SVs, 3.97% were identified as transposition events. However, the list included numerous duplicates or very short variants that were subsequently filtered out. Consequently, we find that after filtering, 7.37% of all SVs are caused by transposition events.

Most of the transposition events were observed due to deletions (60%) while insertions, duplications and inversions cause the remaining variation (File F6 + F7). One difficulty in interpreting these proportions stems from the known biases of sequencing data [102] which make insertions hard to detect. This results in an elevated number of observations of cut transpositions (deletions), but fewer paste transpositions (insertions). Nonetheless, we find certain classes of transposons to be especially active in the comparisons of probe and reference genomes, such as Helitrons and SINEs relative to *VC2010*, and LINEs and Novosib when compared to *CB4856* (Fig. 6 D, File F8). The activity of Helitrons was observed previously [92, 93]. Helitrons were implicated in the divergence of GPCR genes and heat shock elements. Moreover, they are considered to play an important role in evolution [42]. Comparing the two major classes, we conclude that the biggest contribution stems from DNA transposons (82% for *VC2010* comparisons and 95% for *CB4856* comparisons), similar to the findings in [103].

Moreover, we observe a linear relationship between the number of transposition events found and the phylogenetic distance of the given strains (Fig. 6 E-F, Table S9). The strains *QX1211* and *ECA36* have the largest differences based on transposon data before [80].

4 Discussion

Here we present TransposonUltimate, a bundle of three modules for transposon classification, annotation and transposition event detection. Moreover, we present TransposonDB, a database containing more than 891,051 transposon sequences from a wide range of species. Our benchmarks shows that the classification module RFSB outperforms existing methods. Although *RFSB* has a very high accuracy, we believe that performance could be improved by developing species specific classifiers. It would also be helpful to explore new feature representations that strongly correlate to phylogenetic distance metrics.

The annotation module combines existing annotation approaches using an ensemble strategy, and this ensures a less biased outcome than existing methods that tend to favor certain TE classes. The annotation module could be extended by the search for fragmented copies of annotated transposons connected with filters to avoid false positives. Application to three different species revealed that TEs from the same family vary drastically in length. Thus, an important question for future research is to determine to what extent such differences reflect hitherto uncharacterized families, and to what extent the differences correspond to overall sequence divergence.

The detection module enables the identification of transposition events through structural variants in genomes profiled using long-read sequencing technologies. Application of the *deTEct* pipeline to 20 wild type strains of *Caenorhabditis elegans* suggests that transposon events are responsible for 7.37% of structural variants. Although previous studies have argued that transposons are a major driver of structural variation [102], our results suggest that at least for wild isolates of *Caenorhabditis elegans* this is not the case. As additional high quality assemblies become available, it will be interesting to further explore this important question. Moreover, the development of localisation algorithms of target and donor sites of transposons seems a promising add-on for the detection module. Besides, structural variants gathered from whole genome comparison using anchor filtering [104] could be included and compared.

As long-read technologies are becoming more widely used and the number of sequenced genomes rises quickly, there is an urgent need for methods to identify and annotate TEs which correspond to plurality and in some cases a majority of genome sequences. In particular, as more human [105] and other vertebrate (<https://vertebrategenomesproject.org/>) genomes are profiled using these technologies, TransposonUltimate will be a valuable tool to improve our understanding of the impact of TEs on both traits and diseases.

5 Conclusions

Our TransposonUltimate bundle of software tools provides a powerful and user-friendly means of analyzing TEs. In addition to providing highly accurate classifications, our analysis also provides insights as to what features are most informative for predicting TE class. Our ensemble approach to annotation is more unbiased than existing methods that tend to focus on one or a few classes. Finally, our transposition event detection module can take advantage of long-read technologies to identify to what extent TEs underlie SVs.

Competing interests

The authors declare that they have no competing interests.

Author's contributions

The study was conceived and designed by KR, CR, EAM and MH. The code was written by KR, and the analyses were carried out by KR and CR. The work was supervised by EAM and MH. KR and MH wrote the manuscript with input from EAM and CR.

Acknowledgements

We would like to thank Sarah Buddle, Simone Procaccia, Fu Xiang Quah and Alexandra Dallaire for assistance with testing and debugging the software.

Funding

This work was supported by Cancer Research UK (C13474/A18583, C6946/A14492) and the Wellcome Trust (219475/Z/19/Z, 092096/Z/10/Z) to EAM. For the purpose of Open Access, the author has applied a CC BY public copyright licence to any Author Accepted Manuscript version arising from this submission.

Author details

¹Gurdon Institute, University of Cambridge, CB2 1QN Cambridge, UK. ²Wellcome Sanger Institute, Wellcome Genome Campus, CB10 1SA Hinxton, UK. ³Department of Genetics, University of Cambridge, Downing Street, CB2 3EH Cambridge, UK. ⁴Evergrande Center for Immunologic Diseases, Harvard Medical School and Brigham and Women's Hospital, 75 Francis Street, MA 02215 Boston, MA, USA.

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Figures

Tables

Table 1 Overview of common transposon annotation tools. The most commonly used tools such as RepeatMasker and RepeatModeler cover a variety of transposons, while others focus on certain classes only. The tools use one or more of the *de novo*, structural and similarity-based transposon annotation approaches.

Name	Approach			Class I			Class II		
	Novo.	Struc.	Simil.	LTR	LINE	SINE	TIR	HEL	MITE
RepeatMasker	x		x	x	x	x	x	x	x
RepeatModeler	x			x	x	x	x	x	x
CLARI-TE [107]	x	x	x	x	x	x	x	x	x
TESeeker [41]			x	x	x	x	x	x	x
PILER [40]	x			x	x	x	x	x	x
Censor [108]	x			x	x	x	x	x	x
RepLong [109]	x			x	x	x	x	x	x
EDTA [44]	x	x	x	x	x	x	x	x	x
MGEScan [110]	x	x	x	x	x	x			
LTR-Finder [111]		x		x					
LtrDetector [112]		x		x					
LTRpred [73]	x	x	x	x					
LTRharvest [66]	x	x	x	x					
LTRdigest [113]		x		x					
SINE-Finder [68]	x	x				x			
SINE-Scan [69]	x	x				x			
TIRvish [67]		x					x		
HelitronScanner [42]		x						x	
MUSTv2 [70]		x							x
MiteFinderII [71]		x							x
MITE-Tracker [72]		x							x
detectMITE [45]		x							x
MITE-Hunter [47]		x							x

Additional Files

File F1 : TransposonDB.fasta

File F2 : NCBICDD1000.Proteins.txt

File F3 : Classification_FeatureImportanceAnalysis.csv

File F4 : GFF3 files in "PaperSupplements/Annotation/..."

File F5 : GFF3 files in "PaperSupplements/Detection/..."

File F6 : Detection_SVDistribution.csv

File F7 : Detection_PipelineData.csv

File F8 : Detection_ClassDistribution.csv

Supplements

database	before filtering	#F0 no label	#F1.1 fragments	#F1.2 contigs	#F1.3 satellites	#F1.4 RNA	#F1 all #F1.*	#F2 len.100bp	#F3 alphabet	#F4 all rules
ConTEdb	322,705	322,702	322,702	322,702	322,702	322,702	322,702	317,109	317,995	312,402
DPTedB	31,340	31,326	31,326	31,326	31,326	31,326	31,326	31,285	31,325	31,284
PGSB	61,730	61,729	61,729	61,729	61,529	60,813	55,616	60,757	61,397	54,488
MnTEdb	5,925	5,912	5,912	5,912	5,912	5,912	5,912	5,902	5,912	5,902
PMITEdb	2,449,127	357,134	313,211	313,211	313,211	313,211	313,211	298,777	302,529	288,104
RepBase	56,403	53,880	51,674	51,674	51,674	51,648	51,674	51,455	37,796	37,563
RiTE	265,549	264,022	242,216	156,195	241,847	240,509	154,232	152,746	154,058	152,575
SoyetEdB	38,664	38,603	38,603	38,603	38,603	38,603	38,603	38,519	38,601	38,517
SPTEDdb	18,413	18,408	18,408	18,408	18,408	18,408	18,408	18,402	18,408	18,402
TrepDB	4,162	3,910	1,874	3,910	3,910	3,910	1,870	3,669	3,869	1,822
TransposonDB										891,051

Table S1 Classification: TransposonDB filter rule application. This table shows the number or remaining sequences in the constituents databases of TransposonDB after the application of the different filter rules.

(A) Proposed transposon classification taxonomy						
Class	Taxonomy / hierarchy			Notes on constituents		
1	Class I (Retrotransposons)					
1/1	LTR			Copia, Gypsy, Bel-Pao, Retrovirus, ERV		
1/1/1				Copia		
1/1/2				Gypsy		
1/1/3				ERV		
1/2	Non-LTR			DIRs (VIPER, Ngaro), LINEs, SINEs		
1/2/1				R2, RTE, Jockey, L1, I, Randl, Penelope, DRE...		
1/2/2				SINE		
2	Class II (DNA transposons)			TIR, Crypton, Helitron, Maverick / Polinton, MITEs		
2/1	TIR			Tc1-Mariner, hAT, Mutator, Merlin...		
2/1/1				Tc1-Mariner		
2/1/2				hAT		
2/1/3				CMC		
2/1/4				Sola		
2/1/5				Zator		
2/1/6				Novosib		
2/2	Helitron			Helitron		
2/3	MITEs			Tourist, Stowaway		
<div><div>1</div><div>R2 (CRE, R4, Hero, NeSL, R2), RTE (RTETP, Proto2, RTEX, RTE), Jockey (Rex1, CR1, L2, L2A, L2B, Daphne, Crack), L1 (Proto1, Tx1), I (Ingi, Nimb, Tad1, Loa, R1), Randl, Penelope, DRE</div></div>						
<div><div>2</div><div>Tc1-Mariner, hAT, Mutator, Merlin, Transib, P, PiggyBac, PIF-Harbinger, CACTA / ENSPM / Chapaev, MuLE / MUDR, CMC, Sola, Ginger, Academ, Dada, Kolobok, Zator, Novosib</div></div>						

(B) Transposon structure overview						
Structural Features	Class I			Class II		
	LTR	LINE	SINE	TIR	HEL	MITE
Target site duplication (TSD)	x	x	x	x		x
Terminal inverted repeat (TIR)				x		x
Long terminal repeat (LTR)	x					
Primer binding site (PBS)	x					
Polypurine tract (PPT)	x					
Begin A-TC					x	
End CTRR-T					x	
Open reading frames (ORF)	x	x		x		x
Palindromic sequence (hairpin loop)					(x)	
Poly(A) tail		x	x			
Protein Features						
Helicase					(x)	
Capsid protein (GAG)	x					
RPA-like (RAPI) replication protein					(x)	
Envelope (ENV)	(x)					
Transposase				x		
Endonuclease		(x)				
Nucleic acid binding protein (NABP)		x				
Aspartic proteinase (AP)	x					
Apurinic endonuclease (AE)		(x)	(x)			
Pol gene (pol)	x					
Protease (PR)	x					
Integrase (INT)	x					
Reverse transcriptase (RT)	x	(x)				
RnaseH (RH)	x					

Figure 1 Transposon taxonomy and transposon structure. (A) The taxonomy used in this study is based on multiple classification schemes [49], [36], [106], [3] and the taxonomies used by the transposon databases. (B) Autonomous, transposition competent transposons have characteristic structural and protein features depending on their class. The proteins are necessary for the transposons to move via class-specific transposition mechanisms. The x mark which structural and protein features are characteristic to different transposon classes and sub classes for complete, autonomous transposons. The (x) mark features that are not required but if present are indicative.

Figure 4 Evaluation of the RFSB classifier. (A) Benchmark of different transposon classifiers by reported numbers in publications. (B) The performance of selected, reproducible classifiers applied on RepBase+PGSB database using the taxonomy in Fig. 1 A. Reported numbers represent performances from a total perspective. (C) RFSB classification performances from total, taxonomic level and class perspective. (D) Analysis of each feature's contribution to classifier's explanatory power. The white line shows the cumulative explanatory power. (E) Analysis of different k-mer features in combination with protein features for a binary classifier differentiating between class 1 and 2 transposons. All values presented were calculated as average across a tenfold cross validation.

databases	ConTEdb	DPTEdb	mipsREdat-PGSB	MinTEdb	PMTEdb	RepBase23.08	RiTE	Soyetedb	SPTEDdb	TrepDB
Eukaryota	312,334	29,452	52,691	3,154	282,500	36,681	129,911	36,429	6,733	1,068
Animalia	0	0	0	0	0	21,854	0	0	0	20
Chromista	0	0	0	0	256	988	0	0	0	74
Fungi	0	0	0	0	0	1,961	0	0	0	354
Plantae	312,334	29,452	52,691	3,154	282,244	11,721	129,911	36,429	6,733	620
Protozoa	0	0	0	0	0	150	0	0	0	0
UnicellularFlagellate	0	0	0	0	0	7	0	0	0	0
Prokaryota	0	0	0	0	0	54	0	0	0	6
Virus	0	0	0	0	0	31	0	0	0	0
Total	312,334	29,452	52,691	3,154	282,500	36,766	129,911	36,429	6,733	1,074

Table S2 Classification: TransposonDB sequences across biological domains. This table shows the number of sequences in TransposonDB by the source database and biological domains.

databases	ConTEdb	DPTeddb	mipsREdat-PGSB	MnTEdb	PMITEdb	RepBase23.08	RITE	Soyetedb	SPTEdb	TrepDB
Animalia	0	0	0	0	0	21,854	0	0	0	20
Annelida	0	0	0	0	0	287	0	0	0	0
Anthropoda	0	0	0	0	0	8,030	0	0	0	5
Antropoda	0	0	0	0	0	1	0	0	0	0
Ascomycota	0	0	0	0	0	4	0	0	0	0
Bilateria	0	0	0	0	0	1	0	0	0	0
Brachiopoda	0	0	0	0	0	1	0	0	0	0
Chordata	0	0	0	0	0	10,182	0	0	0	11
Cnidaria	0	0	0	0	0	1,228	0	0	0	0
Ctenophora	0	0	0	0	0	31	0	0	0	0
Deuterostomia	0	0	0	0	0	26	0	0	0	0
Echinodermata	0	0	0	0	0	221	0	0	0	0
Hemichordata	0	0	0	0	0	37	0	0	0	0
Mollusca	0	0	0	0	0	749	0	0	0	0
Nematoda	0	0	0	0	0	574	0	0	0	4
Placozoa	0	0	0	0	0	8	0	0	0	0
Platyhelminthes	0	0	0	0	0	343	0	0	0	0
Porifera	0	0	0	0	0	7	0	0	0	0
Priapulida	0	0	0	0	0	1	0	0	0	0
Rotifera	0	0	0	0	0	123	0	0	0	0
Chromista	0	0	0	0	256	988	0	0	0	74
Ciliophora	0	0	0	0	0	11	0	0	0	0
Cliliophora	0	0	0	0	0	3	0	0	0	0
Haptophyta	0	0	0	0	0	15	0	0	0	0
Myxozoa	0	0	0	0	0	104	0	0	0	7
Ochrophyta	0	0	0	0	0	92	0	0	0	0
Oomycota	0	0	0	0	256	760	0	0	0	40
Orchophyta	0	0	0	0	0	3	0	0	0	0
Fungi	0	0	0	0	0	1,961	0	0	0	354
Ascomycota	0	0	0	0	0	749	0	0	0	354
Basidiomycota	0	0	0	0	0	1,111	0	0	0	0
Blastocladiomycota	0	0	0	0	0	14	0	0	0	0
Chytridiomycota	0	0	0	0	0	15	0	0	0	0
Mucoromyceta	0	0	0	0	0	9	0	0	0	0
Mucoromycota	0	0	0	0	0	71	0	0	0	0
Plantae	312,334	29,452	52,691	3,154	282,244	11,721	129,911	36,429	6,733	620
Angiosperms	0	29,452	48,583	3,154	257,898	9,997	129,911	36,429	6,733	618
Bryophyta	0	0	1,060	0	0	45	0	0	0	2
Chlorophyta	0	0	1	0	105	106	0	0	0	0
Rhodophyta	0	0	0	0	0	595	0	0	0	0
Tracheophyta	312,334	0	3,047	0	24,241	978	0	0	0	0
Protozoa	0	0	0	0	0	150	0	0	0	0
Amoebozoa	0	0	0	0	0	78	0	0	0	0
Eozoa	0	0	0	0	0	2	0	0	0	0
Euglenozoa	0	0	0	0	0	14	0	0	0	0
Metamonada	0	0	0	0	0	42	0	0	0	0
Percolozoa	0	0	0	0	0	14	0	0	0	0

Table S3 Classification: TransposonDB sequences across eukaryotic kingdoms. This table shows the number of sequences in TransposonDB by the source database and biological kingdoms.

database	TransposonDB	ConTEdb	DPTedB	mipsREdat-PGSB	MnTEdb	PMITEdb	RepBase23.08	RITE	Soyetedb	SPTEDdb	TrepDB
1	412,975	252,236	23,832	49,853	1,280	0	25,908	23,508	30,538	4,972	848
1.1	360,618	210,831	22,343	48,186	1,267	0	21,542	20,670	30,404	4,769	606
1.1.1	122,715	81,329	5,010	11,059	600	0	6,423	4,277	12,482	1,316	219
1.1.2	144,221	70,605	9,694	18,497	430	0	9,218	14,490	17,922	3,026	339
1.1.3	8,118	4,087	651	0	0	0	3,276	56	0	48	0
1.2	51,943	41,405	1,489	1,253	13	0	4,366	2,838	134	203	242
1.2.1	47,328	40,679	1,440	931	13	0	3,418	272	134	199	242
1.2.2	3,640	0	0	322	0	0	753	2,565	0	0	0
2	478,070	60,098	5,620	2,838	1,874	282,500	10,859	106,403	5,891	1,761	226
2.1	397,819	131	468	1,416	1,752	282,500	7,791	97,710	5,809	59	183
2.1.1	92,563	5	41	261	0	86,195	2,078	2,301	1,645	7	30
2.1.2	94,929	5	144	228	135	0	918	93,494	0	5	0
2.1.3	19,335	25	108	77	1,084	15,076	2,376	487	65	22	15
2.1.4	35,981	0	0	0	0	33,537	0	13	2,370	0	61
2.1.5	145,791	4	93	139	285	142,679	755	131	1,664	8	33
2.1.6	7,552	0	0	711	0	4,996	467	1,280	65	0	33
2.2	56,786	47,199	4,652	14	4	0	665	2,458	82	1,669	43
2.3	20,129	12,767	500	475	118	0	2	6,235	0	32	0
Total	891,045	312,334	29,452	52,691	3,154	282,500	36,767	129,911	36,429	6,733	1,074

Table S4 Classification: TransposonDB sequences across transposon classes. This table shows the number of sequences in TransposonDB by the source database and transposon classes.

Constituents	NCBI CDD ID
Aspartic proteinase	cd00303, cd05481, cd05484
Apurinic endonuclease	tigr00587
Integrase (core domain)	pfam00665, cog3335, pfam13358, pfam01359
GAG pre-integrase	pfam13976
Ribosomal-processing	
cysteine proteinase	cd16332, prk14553
Cysteine proteinase	tigr01586
Peptidase (Prp)	pfam04327
	pfam04231, cog4636, pfam05685, cog2356, pfam01844, pfam05551,
Endonuclease	pfam07510, pfam13391, pfam13392, pfam13395, pfam14414,
	prk15137, cd00719, smart00478, cog0648, prk01060, smart00518,
	prk02308, pfam04493, pfam08459
GAG capsid protein	pfam03732, pfam16297
	smart00490, smart00487, smart00488, smart00491, cog1201,
Helicase	prk13767, tigr04121, pfam06733, pfam00270, pfam00271, pfam04851,
	pfam05970, pfam14617, pfam13307
DNA polymerase	cd08637, cd08638, cd08639, cd08640, cd08641, cd08642, cd08643
RNAse H	cd06266, cd09272, cd09273, cd09274, cd09275, cd09276, cd09279
	prk06863, prk06751, prk06752, prk08182, prk06293, prk06461,
	prk06958, prk07274, prk10053, smart00976, cog0629, cog2965,
Replication protein A	cog3111, prk05733, cog4085, cog3390, prk06341, prk09010,
	pfam00436, tigr00621, pfam02765, pfam04057, pfam08646,
	pfam16686, pfam09104, pfam09103, pfam08661, pfam16900,
	pfam13742
	pfam13966, pfam07727, pfam00078, pfam11474, tigr04416,
Reverse transcriptase	pfam13655, pfam17984, pfam17919, pfam17917, pfam13456,
	pfam06817, pfam06815, cog3344, cd03715, cd03714, cd03487,
	cd01709, cd01699, cd01651, cd01650, cd01648, cd01647, cd01646,
	cd01645, cd01644, cd05471
	nf033179, pfam13006, pfam14706, pfam02281, pfam13701,
	pfam13007, pfam13005, pfam04986, pfam03050, pfam01610,
Transposase (incl. DDE domain)	pfam01609, pfam01548, pfam01526, pfam18759, pfam18758,
	pfam17906, pfam13751, pfam13612, cd01187, cd01186, pfam11427,
	pfam02371, pfam01797, pfam1373, pfam13586, pfam13359, pfam13808,
	pfam13613
Tyrosine recombinase	cd01196, cd01195, cd01194, cd01192, cd01191, cd01184, cd01188,
	tigr02224, prk02436, prk00283, cd00796
Others	cd00397, cd00799, cd06094, pfam03564, pfam05380, pfam05585,
	pfam08284, pfam13975, pfam14223, pfam14244, pfam03184

Table S5 Classification: selection of protein domains This table lists the selected NCBI CDD PSSM model IDs considered for the protein features used in the classification module.

Source	Species	Seq. Technology	#Sequences	Length (BP)	Strain name	Strain location
IRGSP	<i>Oryza sativa subsp. japo.</i>	Illumina	12	374,471,240	Nipponbare	Japan
Wormbase WS279	<i>Caenorhabditis elegans</i>	III., PacBio, Nano.	7	102,092,263	VC2010	Bristol (UK)
Wormbase WS279	<i>Caenorhabditis elegans</i>	Sanger	7	100,286,401	N2	Bristol (UK)
Wormbase WS279	<i>Caenorhabditis elegans</i>	Illumina	7	98,291,416	CB4856	Hawai (USA)
Cristian Riccio	<i>Caenorhabditis elegans</i>	PacBio, Illumina	6	105,149,162	SX3383	Ulupalakua (USA)
Cristian Riccio	<i>Caenorhabditis elegans</i>	PacBio, Illumina	6	103,998,030	SX3399	Wuhan (China)
Cristian Riccio	<i>Caenorhabditis elegans</i>	PacBio, Illumina	6	104,190,248	SX3356	San Francisco (USA)
Cristian Riccio	<i>Caenorhabditis elegans</i>	PacBio, Illumina	6	103,085,866	SX3351	Addisababa (Ethiopia)
Cristian Riccio	<i>Caenorhabditis elegans</i>	PacBio, Illumina	6	103,400,028	SX3368	Adelaide (Australia)
Cristian Riccio	<i>Caenorhabditis elegans</i>	PacBio, Illumina	6	102,737,178	SX3287	Altadena (USA)
Cristian Riccio	<i>Caenorhabditis elegans</i>	PacBio, Illumina	6	102,467,336	SX3304	Amares (Portugal)
Cristian Riccio	<i>Caenorhabditis elegans</i>	PacBio, Illumina	6	104,412,447	SX3306	Auckland (New Zealand)
Cristian Riccio	<i>Caenorhabditis elegans</i>	PacBio, Illumina	6	102,638,870	SX3254	Bristol (UK)
Cristian Riccio	<i>Caenorhabditis elegans</i>	PacBio, Illumina	6	102,222,210	SX3292	Hawai (USA)
Cristian Riccio	<i>Caenorhabditis elegans</i>	PacBio, Illumina	6	102,808,004	SX3302	Hermanville (France)
Cristian Riccio	<i>Caenorhabditis elegans</i>	PacBio, Illumina	6	102,945,196	SX3303	Lake Forest Park (USA)
Cristian Riccio	<i>Caenorhabditis elegans</i>	PacBio, Illumina	6	103,095,877	SX3298	Lisbon (Portugal)
Cristian Riccio	<i>Caenorhabditis elegans</i>	PacBio, Illumina	6	102,627,405	SX3295	Madagascar (Madagascar)
Cristian Riccio	<i>Caenorhabditis elegans</i>	PacBio, Illumina	6	104,465,563	SX3367	Manuka (USA)
Cristian Riccio	<i>Caenorhabditis elegans</i>	PacBio, Illumina	6	103,159,706	SX3305	Palo Alto (USA)
Cristian Riccio	<i>Caenorhabditis elegans</i>	PacBio, Illumina	6	102,930,261	SX3296	Roxel (Germany)
Cristian Riccio	<i>Caenorhabditis elegans</i>	PacBio, Illumina	6	102,990,752	SX3286	Salt Lake City (USA)
Cristian Riccio	<i>Caenorhabditis elegans</i>	PacBio, Illumina	6	103,010,394	SX3350	Southampton (USA)
Cristian Riccio	<i>Caenorhabditis elegans</i>	PacBio, Illumina	6	103,231,504	SX3341	Le Perreux-sur-Marne (France)
JST-ACCEL (NIBB)	<i>Rhizophagus irregularis</i>	PacBio	210	149,750,837	DAOM197198	Quebec (Canada)

Table S6 Annotation: case study genomes. This diagram shows source, species, sequencing technology, number of sequences, length in bp, strain name and location for the 25 case study genomes.

(A) Benchmark of classifiers, performances reported in publications of classifiers (different databases and taxonomies)									
	Random Even	Random Gaussian	HC_GA	HC_LGA	NLLCPN	TERL	TopDown	RFSB	(TransposonDB) RFSB
Overall									
F1	31.03 % (0.32%)	30.44% (0.26%)	83.00%	84.00%	90.00%	85.80%	83.00%	91.34% (0.33%)	89.66% (0.06%)
(B) Benchmark of classifiers, performances reproduced on RepBase+PGSB database and proposed taxonomy									
	Random Even	Random Gaussian	HC_GA	HC_LGA	NLLCPN	TERL	TopDown	RFSB	(TransposonDB) RFSB
Overall									
ACC	47.58% (0.27%)	47.39% (0.38%)	84.19% (0.41%)	80.94% (0.37%)	79.96% (0.28%)	24.10% (2.91%)	38.60% (23.40%)	93.75% (0.24%)	93.28% (0.04%)
F1	31.03 % (0.32%)	30.44% (0.26%)	77.91% (0.59%)	74.02% (0.56%)	70.84% (0.38%)	7.11% (0.99%)	24.43% (24.43%)	91.34% (0.33%)	89.66% (0.06%)
MCC	-11.14% (0.54%)	-11.71% (0.63%)	65.99% (0.86%)	59.45% (0.79%)	57.52% (0.60%)	-54.45% (5.41%)	-27.83% (45.19%)	86.45% (0.52%)	84.68% (0.09%)
Level 1									
ACC	79.23% (0.09%)	79.52% (0.07%)	92.86% (0.13%)	91.78% (0.27%)	92.32% (0.10%)	82.18% (0.54%)	77.99% (5.75%)	96.48% (0.10%)	96.11% (0.02%)
F1	29.52% (0.3%)	29.14% (0.23%)	75.05% (0.47%)	71.17% (0.51%)	70.19% (0.39%)	0.10% (0.09%)	23.05% (21.99%)	88.30% (0.32%)	87.74% (0.07%)
MCC	17.35% (0.36%)	17.18% (0.26%)	70.92% (0.53%)	66.44% (0.67%)	66.70% (0.45%)	-7.36% (0.64%)	10.31% (25.19%)	86.27% (0.38%)	85.44% (0.09%)
Level 2									
ACC	82.60% (0.09%)	82.91% (0.05%)	93.63% (0.13%)	92.72% (0.26%)	93.26% (0.08%)	89.21% (0.09%)	82.33% (3.92%)	96.99% (0.09%)	96.67% (0.02%)
F1	17.57% (0.38%)	17.13% (0.23%)	68.51% (0.73%)	63.70% (0.70%)	60.18% (0.43%)	00.00% (0.00%)	12.89% (20.23%)	86.30% (0.40%)	85.66% (0.08%)
MCC	7.85% (0.43%)	7.62% (0.24%)	65.12% (0.75%)	59.89% (0.78%)	59.21% (0.48%)	-1.01% (0.50%)	3.21% (22.08%)	84.65% (0.45%)	83.81% (0.09%)
Level 3									
ACC	86.30% (0.05%)	86.57% (0.04%)	92.87% (0.24%)	91.69% (0.22%)	90.91% (0.00%)	90.90% (0.01%)	86.96% (3.16%)	97.33% (0.11%)	97.76% (0.02%)
F1	9.63% (0.41%)	9.32% (0.34%)	49.93% (2.77%)	38.67% (4.62%)	0.00% (0.0%)	0.00% (0.00%)	7.96% (12.97%)	84.95% (0.59%)	87.51% (0.10%)
MCC	2.59% (0.42%)	2.56% (0.34%)	48.61% (2.44%)	37.48% (3.24%)	0.00% (0.00%)	-00.21% (0.17%)	2.28% (12.87%)	83.51% (0.65%)	86.29% (0.11%)

Table S7 Classification: benchmark of classifiers. All performance measures reported as average across 10 folds (in %) are supplemented by the standard deviations in brackets (in %). Bold numbers mark the best performance amongst different classifiers within same category. Panel (A) outlines performance measures of the benchmark algorithms reported in their publications (meaning these results were gathered from different datasets and taxonomies, depending on the specific publication). Panel (B) outlines performance measures of several benchmark algorithms to the proposed "RFSB" classifier methodology. All results were calculated based on the same dataset RepBase+PGSB and the same, proposed taxonomy. The measures are reported for taxonomic levels and overall perspective. In addition, the proposed "RFSB" classifier is applied to TransposonDB and reported in the most right column.

strain	VC2010		CB4856	
	PBSV (pbmm2)	Sniffles (ngmlr)	PBSV (pbmm2)	Sniffles (ngmlr)
SX3383 (QX1791)	665	848	295	289
SX3399 (GXW1)	384	546	278	27
SX3356 (QX1211)	862	831	371	297
SX3351 (DL200)	354	390	260	237
SX3368 (AB1)	262	288	284	233
SX3287 (PS2025)	521	571	257	210
SX3304 (EG4725)	528	417	268	185
SX3306 (ECA36)	927	837	372	299
SX3254 (N2)	13	10	246	212
SX3292 (CB4856)	670	744	2	28
SX3302 (JU394)	208	252	273	234
SX3303 (JT11398)	275	309	262	220
SX3298 (JU775)	494	509	271	55
SX3295 (LKC34)	378	336	261	203
SX3367 (DL238)	671	861	279	287
SX3305 (ECA248)	454	411	271	225
SX3296 (MY23)	555	472	279	187
SX3286 (EG4946)	277	272	270	214
SX3350 (PB306)	458	540	284	258
SX3341 (JU751)	431	424	263	192

Table S8 Detection: Number of observed transposition events. This table shows the number of observed transposition events for different probe reference genome combinations, alignment and structural variant calling tools.

strain	Number of transposition events										Phylogenetic distance	
	Sniffles-NGMLR					PBSV-PBMM2						
	Total VC2010	CB4856	TotalBP VC2010	CB4856	Total VC2010	CB4856	TotalBP VC2010	CB4856	TotalBP VC2010	CB4856	VC2010	CB4856
AB1	288	233	2528558	864269	262	284	450579	907524	0.023819	0.094459		
CB4856	744	28	3508318	86028	670	2	1350133	11185	0.096336	0		
DL200	390	237	1260330	918529	354	260	737484	613060	0.078775	0.144549		
DL238	861	287	5340946	1120948	671	279	1635629	701238	0.100631	0.077201		
ECA248	411	225	1674096	828618	454	271	861328	978863				
ECA36	837	299	5547803	933204	927	372	2241605	779615	0.193356	0.185248		
EG4725	417	185	1116717	976099	528	268	1077970	542523	0.063796	0.104196		
EG4946	272	214	948485	1126127	277	270	839962	797286	0.036709	0.093291		
GXW1	546	27	1990299	42255	384	278	784283	624929	0.036727	0.096107		
JT11398	309	220	866883	1137900	275	262	557395	664228	0.027652	0.093426		
JU394	252	234	1413635	1200411	208	273	747156	795337	0.014571	0.109029		
JU751	424	192	4025712	880873	431	263	971156	842471	0.043152	0.099734		
JU775	509	55	2324057	198857	494	271	1175322	589865	0.115199	0.165997		
LKC34	336	203	935153	763459	378	261	909664	710945	0.037956	0.088754		
MY23	472	187	2079515	480242	555	279	1227010	500555	0.08266	0.139242		
N2	10	212	39344	576541	13	246	20414	520726	0	0.096336		
PB306	540	258	2226567	1042300	458	284	854083	735233	0.056832	0.113414		
PS2025	571	210	3381679	856256	521	257	1018228	738095	0.071363	0.127945		
QX1211	831	297	3556466	936120	862	371	1895254	723441	0.194891	0.186783		
QX1791	848	289	5571466	1232941	665	295	1521737	860000	0.05031	0.097823		

Table S9 Detection: Genetic distance and number of transposition events found. This table shows the number of observed transposition event candidates, the length of their mask in bp, and the phylogenetic distance of probe and reference genome.