DupMasker: A tool for annotating primate segmental duplications

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Segmental duplications (SDs) play an important role in genome rearrangement, evolution, and the copy-number variation (CNV) of primate genomes. Such sequences are difficult to detect, a priori, because they share no defining sequence features that distinguish them from unique portions of the genome. Current sequence annotation of segmental duplications requires computationally intensive, genome-wide self-comparisons that cannot be easily implemented on new data sets. Based on the successful implementation of RepeatMasker, we developed a new genome annotation tool, *DupMasker*. The program uses a library of nonredundant consensus sequences of human segmental duplications, wherein a majority of the ancestral origins have been determined based on comparisons to mammalian outgroup genomes. Using *DupMasker*, new human and nonhuman primate (NHP) sequences may be readily queried to provide details on the origin and degree of sequence identity of each duplicon. This program can be applied to delineate the order and orientation of duplicons within complex duplication blocks and used to characterize structural variation differences between sequenced human haplotypes. We predict this tool will be valuable in the annotation of large-insert sequence clones, allowing putative unique and duplicated regions of the genomes to be annotated prior to whole genome assembly comparisons.

[Supplemental material is available online at www.genome.org.]

Initial analysis of the human genome and other primate genomes reveals that 4%–6% of each genome is composed of segmental duplications (Bailey et al. 2001, 2002; Cheung et al. 2003; Chen et al. 2004; She et al. 2004, 2006; Sainz et al. 2006). We now know that segmental duplications are hot spots for non-allelic homologous recombination (NAHR), copy-number variations (CNVs), and genomic rearrangements, leading to more than two dozen genomic diseases (Lupski 1998). The organization of human segmental duplications is complex. They are arranged into duplication blocks of mosaic architecture made up of many independent duplication events (termed duplicons) that have both shared and independent evolutionary histories (Jiang et al. 2007). These patterns are difficult to discern based solely on pairwise alignments and usually require detailed hand curation to delineate the evolutionary breakpoint boundaries.

Current methods used to detect segmental duplications are based on a self-comparison of the entire genome or based on comparison of whole genome shotgun sequence data against a reference genome (Bailey et al. 2001, 2002). These methods have two notable limitations. First, the existing pipelines are computationally intensive and are not easily implemented on new genome assemblies or incomplete data sets. Second, the output of these available methods provides limited information regarding the substructure, relationship, or ancestral origin of the segmental duplications (Jiang et al. 2007). As a result, cross-comparison between loci or species is limited to a series of pairwise alignments and is complicated by the difficulty of mapping between incompletely sequenced paralogs.

Taking advantage of the consensus sequence library and ancestral state information provided by our previous study (Jiang et al. 2007), we developed the software *DupMasker*, which (1) defines

the orientation of individual duplicons for a given primate genomic sequence, (2) delineates the fine mosaic substructure for a given complex duplication block, and (3) provides information regarding the ancestral origin for 70% of human segmental duplications.

Results

We developed *DupMasker* in three basic steps. We first constructed a library (duplib) of consensus sequences for duplication subunits (size ≥ 100 bp) (Jiang et al. 2007), which captures 97.2% of the sequence information within the human set of segmental duplications ($\geq 90\%$ identity and ≥ 1 kb in length). Previously, we decomposed all human 28,856 pairwise alignments into a nonredundant set of 12,087 duplication subunits using a modified “A-Bruijn” graph algorithm. Of these, the ancestral origin could be determined for 67.3% by comparison with mammalian outgroup species. We generated a representative consensus sequence for each of these duplicons and identified each duplicon by its ancestral map location in the human genome, adding biological definition to the library. We note, however, that the ancestral location for ~30% of the duplicated base pairs (particularly those organized as tandem clusters) is currently impossible to resolve due to gene conversion (a.k.a. concerted evolution) or ambiguous mapping to ancestral mammalian species. In these cases, the duplications are simply represented as human duplication subunit consensus as opposed to ancestral duplicons.

The next step integrates the duplication library into a modified version of RepeatMasker, which performs a sequence comparison of query sequence and consensus sequences within duplib. The procedure initially excludes common primate repeat sequences using RepeatMasker libraries specific for each species. Seed alignments are then generated based on comparing the remaining input sequence to the human duplication library. Next,

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Article published online before print. Article and publication date are at <http://www.genome.org/cgi/doi/10.1101/gr.078477.108>.

Figure 1. *DupMasker* defines the substructure of human segmental duplication blocks. Human segmental duplications are organized into complex duplication blocks where individual duplcons originate from different regions of the genome. We assessed the ability of *DupMasker* to accurately define these ancestral duplcons in this context by comparing results from two regions studied previously in detail (Horvath et al. 2000; Jiang et al. 2007). We schematically display (PARASIGHT: <http://eichlerlab.gs.washington.edu/jeff/parasight/index.html>) duplcons detected using the A-Brujin graph approach (top) versus *DupMasker* (bottom) for (A), an ~600-kbp region on chromosome 2p11 and (B), an ~700-kbp region on chromosome 5q13.2. The different duplcons are illustrated as color-coded blocks; different colors correspond to different cytogenetic band locations of the ancestral loci. We found 33/36 nonredundant duplcons blocks are consistent between these two results. The three mismatched blocks are relatively small in length (length <1.5 kb, highlighted in red).

duplcons are clustered according to ID and consensus agreement, and edges are extended along the query sequence until either a consensus is exhausted or a region of nonrepeat masked

sequence (>7 kb) is encountered. This length boundary was selected as the upper bound for most retrotransposon L1 insertions. The clusters with similar IDs are then grouped, and groups

Table 1. A chromosome comparison of segmental duplications—*DupMasker* vs. WGAC

	<i>DupMasker</i>	WGAC	Shared	Missed	Novel
chr1	12242450	9989848	9321899	667570	2920551
chr2	11430232	9565952	9115241	450407	2314991
chr3	4764816	3139917	2740273	399422	2024543
chr4	6770601	4832059	4314690	517205	2455911
chr5	7370643	5792227	5525762	266243	1844881
chr6	5250037	3378346	3216118	162082	2033919
chr7	14657505	12899369	12151339	747678	2506166
chr8	4295216	2939651	2567565	371928	1727651
chr9	12095511	11638137	10836617	801185	1258894
chr10	10040693	8786118	8269055	516761	1771638
chr11	6909441	5430990	4950296	480500	1959145
chr12	4257900	2841984	2569490	272330	1688410
chr13	3715651	2927545	2695437	231987	1020214
chr14	3622398	2633208	2352963	280134	1269435
chr15	8423129	7967804	7405538	561831	1017591
chr16	8699010	7748086	7350556	397321	1348454
chr17	7682604	6933336	6585851	347229	1096753
chr18	2288338	1867604	1693866	173664	594472
chr19	7841729	3999874	3657240	342488	4184489
chr20	1891551	1438188	1307929	130170	583622
chr21	2079015	1825709	1709678	115967	369337
chr22	4406162	4028612	3632451	396034	773711
chrX	13035873	10314126	9826474	487299	3209399
chrY	13548797	12315095	11560129	754828	1988668
Sum	177319302	145233785	135356457	9872263	41962845
Percentage			0.932	0.068	0.237

We compared the duplication intervals defined by *DupMasker* against those defined by the WGAC method. This table shows the nonredundant base pairs between these two methods. Shared indicates bp consistent between these two methods; Missed, positive by WGAC but negative by *DupMasker*; and Novel, positive by *DupMasker* but negative by WGAC. We also performed a *DupMasker* analysis on build36, and we found there is a slight increase (2.2%) in the amount of segmental duplications between build36 and build35 (181.3 Mb vs. 177.3 Mb). The software package and example files of *DupMasker* can be downloaded at <http://www.repeatmasker.org/DupMaskerDownload.html>.

of consensus and bounded query regions are realigned using WUBLAST2 (Washington University BLAST version 2.0).

The current version of the program uses a human library consisting of 12,087 duplication subunits and generates two standard outputs. These outputs include a file containing the duplication seed information and a second file that contains the information of locally chained duplicons, including duplication subunit ID and orientation in respect to consensus sequence and ancestral locus information.

Based on this design, we constructed a prototype version of *DupMasker* and assessed its efficacy by benchmarking it against previously annotated human segmental duplications mapping to chromosome 2p11 and 5q13.2 regions (Fig. 1A,B). A comparison of previously validated duplication structures with those determined by *DupMasker* shows very good correspondence (33/36 duplication subunits correctly identified with previous annotated sequences) (Horvath et al. 2000; Jiang et al. 2007). Several limitations were noted, especially in the treatment of repeats within or near the boundaries of segmental duplications. For example, some smaller subunits were not identified simply due to overlap with low-complexity repeat sequence. More importantly, the enrichment of common repeats at the boundaries of the duplicons significantly limited our precision in defining the edges of each duplicon using the initial prototype. To eliminate potential repeat-induced artifacts, we excluded all duplicons that contained <50 bp of nonrepeat sequence. Finally, we empirically assessed differential weighting schemes to improve junction detection. Based on these modifications, we estimate that ~93.2% of human input sequence can now be correctly annotated as segmental duplication (Table 1).

In order to assess the validity of *DupMasker* as a stand-alone program to accurately identify segmental duplications, we analyzed the entire human genome (build35) using *DupMasker* and compared the consistency between *DupMasker* results versus Whole Genome Assembly Comparison (WGAC) data (Table 1). Overall, 93.2% of duplications (135.35 Mb/145.23 Mb) are consistent (shared) between these two methods. A relatively small fraction (6.9% or 9.87 Mb) was identified by WGAC but not detected by *DupMasker* as a segmental duplication. Sequence analysis of these “missed” duplications showed that the majority (8.99 Mb/9.87 Mb = 91.1% by base pair composition) corresponded to common repeat sequences. Such losses are expected for segmental duplications enriched in common repeats due to the initial triage design of *DupMasker*, which excludes repeat regions.

In contrast, *DupMasker* predicted 41.96 Mb of duplications that were not

originally classified using the WGAC method. We termed these *DupMasker*-only duplications as “novel” segmental duplications. Similar to RepeatMasker, *DupMasker* has the ability to detect smaller and more divergent duplications (>75% identity with respect to the consensus and less than 1 kbp in length). The WGAC procedure operationally defines segmental duplications as pairwise alignments 1 kbp or more and 90% or more sequence identity. We therefore assessed the length and percentage of identity distribution of these putative “novel” SDs. We found that 91.0% (38.17/41.96 Mb) of these duplications were less than 1 kb (Fig. 2A). More than half of these novel intervals are common repeats (21.95/41.96 Mb) due to the imprecision of boundary definition within repeat-rich regions. We also performed a modified WGAC analysis on the 41.96 Mb using a more relaxed threshold (nonrepetitive sequence alignment size 100 bp or more and BLAST-sequence identity 75% or more) than that of standard WGAC. This modified WGAC analysis identified alignments for 31% (13.1/41.96 Mb) of these “novel” SDs. Among these 13.1 Mb alignments, 97.7% (12.8/13.1 Mb) represent either small (size < 1 kbp) or relatively

Figure 2. The size and sequence identity distribution of “novel” duplications. (A) The length distribution of *DupMasker* duplications not detected by WGAC (termed “novel” SDs) reveals that the majority (99% by number of intervals, 91% by base pair) of these intervals are small fragments (size <1 kb). (B) We found 52.3% (21.9 Mb) of these small intervals are common repeats due to imprecision of boundary definition within repeat-rich regions. We performed a modified WGAC analysis using a relaxed threshold (require nonrepeat alignment ≥ 100 bp and sequence identity $\geq 75\%$) on these “novel” SDs. The analysis revealed alignments for 31% (13.1/41.96 Mb) of these “novel” SDs. Among the 13.1-Mb alignments, 97.7% (12.8/13.1 Mb) represent either small (size <1 kbp) or relatively ancient duplications (sequence identity <90%).

ancient duplications (sequence identity < 90%) (Fig. 2B; Supplemental Table 1).

We tested three different applications of *DupMasker*: the analysis of regions flanking genomic disorders, the analysis of sequence from regions of structural variation, and a genome-wide analysis of a nonhuman primate genome assembly. Results from these various applications illustrate the utility of this software tool.

Analysis of regions associated with genomic disorders

Duplication-rich regions of the human genome are hotspots of NAHR, leading to many human diseases, known as genomic disorders (Lupski 1998; Sharp et al. 2005, 2006, 2008; Mefford et al. 2007). Delineating the duplication architecture of those regions and their underlying LCRs (low copy repeats) or duplcons is important for understanding not only the evolutionary origin but likely sequences that promote non-allelic homologous recombination. *DupMasker* allows the duplication architecture flanking these regions to be decoded and provides information regarding the divergence and orientation of each individual fragment. Figure 3 is a schematic showing the architecture, as predicted by *DupMasker*, one of the most unstable regions of the human genome associated with Prader-Willi syndrome, and a recently described mental retardation syndrome (Sharp et al. 2008). *DupMasker* identifies candidate duplcons of high-sequence identity and proper orientation (color-coded boxes). The duplication architecture corresponds to breakpoints defined by arrayCGH experimental results (highlighted by dashed lines in

Fig. 3). These results highlight the utility of *DupMasker* to predict regions of potential instability associated with NAHR-mediated microdeletion syndromes.

Analysis of sequenced clones

Another application for *DupMasker* is to annotate the duplication composition of sequenced clones, such as fosmid or BAC clones. This can be used to readily exclude certain regions for PCR or oligonucleotide design based on the underlying copy-number and sequence identity of the duplications. Moreover, regions of copy-number variation are particularly enriched in segmental duplications (Sharp et al. 2005; Redon et al. 2006), and annotated duplication maps of two sequences can be used to reconstruct the series of rearrangements that have occurred between any two human haplotypes. Since many of the segmental duplications are shared between humans and other nonhuman primate species, this is particularly valuable when characterizing nonhuman primate sequences that appear rearranged compared with the human genome. Figure 4 shows examples of structural variation between human haplotypes and between species that can be characterized using *DupMasker*. Figure 4A reveals a large deletion in human individual (ABC9) mediated by an NAHR between flanking duplcons, while Figure 4B depicts a lineage-specific segmental duplication insertion event in chimpanzee compared with the corresponding human sequence. We predict that *DupMasker* will be particularly valuable in annotating the breakpoints of CNVs and speciation chromosomes, which are signifi-

Figure 3. Duplication architecture flanking genomic disorders. This figure shows the duplication architecture defined by *DupMasker* for one of the most unstable regions of the human genome (15q11–15q13). (A) Blue lines delineate intrachromosomal duplications of high-sequence identity (size ≥ 10 kb and sequence identity $\geq 95\%$) within this region (WGAC) and identify four breakpoint regions associated with Prader-Willi/Angelman Syndrome and the 15q13.3 deletion syndrome. (B) The duplication substructures defined by *DupMasker* are depicted as color-coded boxes with different colors representing different cytogenetic band locations of duplcons. (C) ArrayCGH data from one patient with Prader-Willi syndrome (bottom) and two patients with chr15q13.3 deletion (Sharp et al. 2008) indicate the patients' deletion breakpoints overlap with the duplcons defined by *DupMasker*. The locations of the breakpoint intervals are highlighted by red dashed lines.

Figure 4. Genomic comparisons by *DupMasker*. *DupMasker* facilitates the characterization of duplication-mediated genomic rearrangements. (A) Miropeats (Parsons 1995) comparison between human reference genome (build35, *top*) against a fosmid clone (*bottom*) from a Japanese individual (ABC9) identifies a ~40-kbp deletion. *DupMasker* on this region identified a pair of tandem duplications (dark green) flanking the internal duplication (light green), which was likely deleted by NAHR in this Japanese individual. The deletion removes part of the intron of the *LATS1* gene. (B) A similar comparison between sequences from a chimpanzee BAC clone (AC097264.4) and its orthologous locus on human chromosome 17 predicts a large (~80 kbp) chimpanzee-specific insertion. *DupMasker* analysis suggests that the insertion is the result of a duplicative transposition event composed of segmental duplications that originated from human-chimpanzee ancestral chromosome 16.

cantly enriched for segmental duplications (Armengol et al. 2003; Bailey et al. 2004).

Analysis of nonhuman primate genomes

Since the consensus sequence library is based on human sequence, it will be necessary to update the library to include species-specific duplications from other nonhuman primate genomes as they are identified. In this regard, *DupMasker* greatly facilitates the identification of lineage-specific duplications. For example, if we apply *DupMasker* (human duplib) to a nonhuman primate genome assembly, we can compare *DupMasker* regions in the NHP genome (duplicated in human) against regions predicted to be duplicated by independent analyses of those genomes (predicted to be duplicated within the NHP by WGAC/WSSD). Such analyses will readily distinguish three types of duplications: duplications shared between human and the NHP, duplications specific to human, and duplications specific to the NHP. Figure 5 illustrates the way different types of duplication (e.g., lineage-specific or shared duplications) can be identified through a comparison of different duplication analyses on the macaque genome (Gibbs et al. 2007) compared with those de-

tested by *DupMasker*. This comparison of the macaque genome predicts that 22.3 Mb are shared duplication between macaque and human, while 122.9 Mb emerged within the human lineage and only 24.3 Mb emerged within the macaque lineage since divergence.

Discussion

We have developed an annotation tool that allows the complex duplication structure of regions to be deciphered and compared without the need for initiating a genome-wide self-comparison. The annotation provides insight into the origin, degree of sequence identity, and orientation of duplicons embedded within sequence. Since many segmental duplications recurrently duplicate (Johnson et al. 2006) or have been shared among species closely related to human, the distribution of this tool will enhance the sequence and assembly of complex regions of great ape genomes by allowing annotators within the sequence centers to distinguish unique from duplicated regions. “DupMasking” of BACs will flag potential regions of new insertion that can then be further characterized. Similar to RepeatMasker, distribution of this tool will have other more pragmatic uses to genetics and genome

Figure 5. Assigning lineage-specific and shared duplications in primates. We applied *DupMasker* (standard default settings) to the macaque genome (RheMac2) and readily identified shared and lineage-specific duplications by comparing the results with duplication maps of the Rhesus Macaque Genome (Gibbs et al. 2007). We found that 84% (121.0/143.3 Mb) of duplications in the human genome are human-lineage specific. There are 22.3 Mb of duplications shared between human and macaque, and 24.3% (24.3/46.4 Mb) of duplications defined in the macaque genome that are macaque-lineage specific.

research, ranging from enhancing oligonucleotide PCR design to improving genotyping assays. Many commercial/customized platforms for SNP genotyping wish to avoid highly duplicated regions of the genome. Our tool not only allows such regions to be identified but also provides information on the copy number of each segment within the reference genome assembly (Supplemental Table 2).

Further enhancements will entail the modification of the duplication library specifically for each nonhuman primate species. We anticipate the discovery of a significant number of lineage-specific duplications (and deletions) in different primate genomes (Cheng et al. 2005). As these regions are discovered, the human duplication library will be modified accordingly to include chimpanzee-specific and macaque-specific duplications. The annotation of BAC sequences will be particularly useful in this regard since we recognize that lineage-specific duplications will occur nonrandomly in the genome (i.e., in the vicinity of shared duplication blocks). Thus, as BAC insert sequences are annotated using WSSD (Bailey et al. 2002) and *DupMasker*, new regions of duplication will be identified. These sequences can be extracted and added to the species-specific duplib as part of the reiterative process of modifying the human duplib. Ultimately, a duplication library specific for each of the primates will emerge.

In addition, we now know that duplication regions are hotspots for extensive copy-number and structural variation. Considering that duplication-mediated NAHR is the most com-

mon mechanism leading to copy-number variation (Kidd et al. 2008), we predict that *DupMasker* will aid in characterizing the duplication architecture of these regions as more copy-number variant regions become sequenced (Fig. 4).

Methods

Duplication library

We developed a library of consensus sequences (duplib) based on the WGAC human segmental duplication data set. The initial data set consisted of 28,856 pairwise alignments (sequence identity $\geq 90\%$ and size ≥ 1 kb) defined by the WGAC method (build35) (Bailey et al. 2001). We applied a modified A-Bruijn graph approach (Pevzner et al. 2004; Jiang et al. 2007) to convert pairwise alignments into nonredundant duplication subunits ($n = 12,087$, size ≥ 100 bp), as described previously. A set of consensus sequences was generated for each duplication by identifying the majority-rule nucleotide within each multiple sequence alignment. The available ancestral state information (102.4 Mb/67.2% of all duplications) for duplication subunits was defined by a reciprocal best-hit between human and outgroup mammalian genomes (Jiang et al. 2007).

DupMasker design

The program initially screens input sequences for all common interspersed repeats using standard RepeatMasker settings (primate library). Repeat-masked base pairs are replaced with Ns, and seed alignments are identified between duplib and the masked test sequence using WUBLAST2 (minimal BLAST score = 300). These seed alignments are stored as part of the *.dupout file. We extend seed alignments by combining local fragments. Local collinear seeds (adjacent seeds from the same duplicon, in the same orientation, and within a default gap length of ≤ 7 kbp) are first chained. Next, the chained query sequence is realigned against the unmasked consensus sequence in the library. The realignment results are stored as part of the *.duplicons output file. The program uses a simple UNIX command line format: segdupmask [-options] [input DNA sequence file]. There are four basic options: (1) -maxDiv restricts the maximal divergence (sequence identity) between the seeds and the consensus sequence; (2) -maxWidth restricts the maximum non-repetitive/nonseed realign gaps (default is 7 kb) for chaining; (3) -forceSearch forces the program to perform all steps despite the presence of previous result files (by default the program will select previous *.dupout and *.out for a given input sequence, omitting the first two steps of the procedure); and (4) -align option generates alignments as part of the standard output. The input file for *DupMasker* is a single text file containing the DNA sequence in FASTA format. After the execution, *DupMasker* creates two standard output files: (1) a text file containing information of all seed alignments (*.dupout) and (2) a text file containing information of all chained duplicons (*.duplicons) with ancestral state information.

Acknowledgments

We thank Pavel Pevzner and Haixu Tang for assistance in implementation of the modified A-Bruijn graph theory algorithm; Heather Mefford, Jeffrey Kidd, and Tonia Brown for useful comments; and Lin Chen for computational assistance. This work was supported by an NIH grant GM058815 to E.E.E. and a Rosetta Inpharmatics fellowship (Merck Laboratories) to Z.J. E.E.E. is an investigator of the Howard Hughes Medical Institute.

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Received March 17, 2008; accepted in revised form May 19, 2008.

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Genome Res. 2008 18: 1362-1368 originally published online May 23, 2008
Access the most recent version at doi:[10.1101/gr.078477.108](https://doi.org/10.1101/gr.078477.108)

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