Oral Spirochetes Implicated in Dental Diseases Are Widespread in Normal Human Subjects and Carry Extremely Diverse Integron Gene Cassettes

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Integrons are genetic elements that acquire and excise gene cassettes from their locus via site-specific recombination. The type of integron first discovered in the 1980s as the source of antibiotic resistance determinants (37) has been named the resistance integron, or the mobile integron, as it is often found in plasmids or associated with transposons. Another type of integron, the chromosomal integron, was discovered in 1998 from examination of the \textit{Vibrio cholerae} genome (22). Although they have similar structures, the two types of integrons (mobile integrons and chromosomal integrons) have different evolutionary histories and differ in that the mobile integrons usually carry relatively few genes, which are predominantly antibiotic resistance genes, while chromosomal integrons often carry far more genes, of very diverse functions (4). Chromosomal integrons are the ancestors of resistance integrons (34).

Integrons consist of a site-specific tyrosine recombinase (\textit{intI}) gene, the primary recombination site \textit{attI} immediately adjacent to the \textit{intI} gene, and an array of captured gene cassettes encoding accessory functions (5). Gene cassettes are the minimal units that can be mobilized by the integrase, with each cassette containing one or a very small number of genes (6), and are separated by a recombination site, \textit{attC}. Aggregation of different gene cassettes results in variable gene cassette arrays. The number of gene cassettes in integrons can reach several hundred; for example, the total length of the gene cassette pool from five \textit{Vibrio} chromosomal integrons is equivalent to a small genome (34).

PCR with degenerate primers targeting the conserved regions of \textit{attC} sites has recovered novel integrase genes and hundreds of diverse gene cassettes from various environments, including soil, sediment, biomass, and water habitats (24, 33, 38). Rowe-Magnus et al. employed a three-plasmid genetic strategy to recover integron genes, using the integrase to bind integron \textit{attC} sites (34). These methods, which utilized the conserved nature of integron recombination sites, revealed a very dynamic integron gene repertoires and suggested that the gene cassette pool is likely to be limitless (7), while at the same time we do not know of work identifying the sources of integron genes.

Here, we report a different approach to discovering chromosomal integrons in human-associated microbial communities, using shotgun metagenomic sequences of the human microbiomes. Human bodies are complex ecological systems, in which various microbial organisms and viruses interact with each other and with human hosts. The MetaHit project has established a human gut microbial gene catalogue (29) and defined three enterotypes of human gut microbiomes (2). The Human Microbiome Project (HMP) (25) has resulted in >700 data sets of shotgun metagenomic sequence (http://www.hmpdacc.org/), from which we can learn the compositions and functions of human-associated microbial communities.

Our approach to integron discovery builds upon two novel computational methods: a targeted assembly approach for identifying the \textit{attC} sites associated with chromosomal integrons (the repeats) in reads and a constrained assembly approach for identifying the gene cassettes, which first avidly retrieves potential paths in the de Bruijn graph (8, 26) for a metagenomic data set, constrained to contigs containing the \textit{attC} sites, and then selects the paths that most likely represent cassette genes, based on several parameters.
As we show in Results, such specialized computational tools are important for a comprehensive characterization of metagenomic functional elements that contain repeats (such as the \textit{attC} sites in the integron gene casettes), as these repetitive regions are extremely difficult to assemble using a whole-metagenome assembly strategy.

In this study, we focus on the identification and characterization of integrons associated with \textit{Treponema} species implicated in periodontal disease (9,36) in the HMP data sets, using our integron discovery system. The \textit{Treponema denticola} genome contains a chromosomal integron with 45 gene cassettes (6), and it is the only human-associated bacterial species that harbors chromosomal integrons (5). We also discover that the draft assemblies of two HMP reference genomes, \textit{Treponema vincentii} and \textit{T. phagedenis}, contain integron \textit{attC} sites similar to those in \textit{T. denticola} and possibly harbor integrons. We do not find integrons in other \textit{Treponema} species, including \textit{T. pallidum} SS14 uid58977 (21), \textit{T. pallidum} Nichols uid57585 (12), \textit{T. primitia} ZAS-2, and \textit{T. azotonutricium} ZAS-9 (13). From the HMP data sets, we identify 826 integron gene cassettes that are related to the \textit{Treponema} species, providing a gene cassette pool with 598 nonredundant genes. With these newly identified gene cassettes, we are able to compare the gene cassettes from different human subjects and study the dynamics of the integron gene cassettes in their natural environments (i.e., human bodies), providing a first survey of integron-containing \textit{Treponema} species and their integrons in a normal human population.

**MATERIALS AND METHODS**

Choosing representative repeat sequences for \textit{Treponema} chromosomal integrons. Eight distinct sequences were selected to represent the integron \textit{attC} sites in the \textit{T. denticola} genome (the complete genomes of the other two integron-containing \textit{Treponema} species, \textit{T. vincentii} and \textit{T. phagedenis}, are not available), given that not all the repeats are identical (Fig. 1). The pairwise sequence similarity between these eight sequences ranges from 77% to 44%, and all the \textit{attC} sites in \textit{T. denticola} can be aligned with at least one of the representative sequences with 85% sequence identity. Once the representative sequences are selected, we are able to identify new \textit{attC} sites using similarity searches, instead of looking for features of integron recombination sites as in reference 39. One advantage of using similarity searches is that we can recover degenerate sites that may lack some typical characteristics of integron recombination sites.

Targeted assembly to identify integron \textit{attC} sites. The targeted assembly approach was developed to characterize CRISPR arrays from shotgun metagenomic sequences (31) and was employed here to identify and assemble the integron \textit{attC} sites. The steps were (i) searching for reads that contain \textit{attC} sites (with identity of >70% and covering >50% of at least one of the representative \textit{attC} sequences) using BLAST (1); for paired-end reads, if one of a pair qualifies, both reads for the pair are included; and (ii) assembling the retrieved short reads using SOAPdenovo (18); note that we used \textit{k}-mers of 31 bp, which were sufficiently long to assemble reads with

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**FIG 1** (A) The neighbor-joining tree of the eight representative sequences of the \textit{T. denticola} chromosomal integron recombination sites. The sequences are named by the starting position of the sites in the genome. The multiple alignment was prepared using ClustalW, and the neighbor-joining tree was prepared using the jalview tool. (B) Predicted structure of one of the representative sequences, \textit{attC1870410}, which has the typical structure of an integron recombination site, with two stems and one conserved unpaired G. The structure was predicted by RNAscf (3), software that performs simultaneous alignment and folding of RNAs, using the eight representative sequences as input.
the repetitive sequences found in the integrons; by contrast, whole-metagenome assembly generally uses shorter k-mers (29).

**Constrained assembly to retrieve integron gene cassettes.** A second method, constrained assembly, was used to assemble integron gene cassettes from metagenomic shotgun reads. Since integron cassettes consist of genes that are much longer than the read length (~100 bp for the current Illumina technology), and the attC sites behave like repeats that confuse (meta-)genome assemblers, it is extremely difficult to obtain gene sequences using either a whole-genome assembly method or the targeted-assembly approach (which is good for assembly of repeats but does not assemble very far beyond the repeats). As the integron cassettes are bounded by two attC repeats, we took advantage of this structure and devised a novel way to retrieve the cassette genes by traversing the assembly graph, constrained by the edges (contigs) that contain the attC sites. To avoid introducing artificial integron genes, we further applied several criteria to select paths that are most likely to present genuine gene cassettes. The constrained assembly approach consists of the following steps (Fig. 2): (i) assembling all shotgun reads in a metagenomic sequence data set—along with the contigs constructed by the targeted assembly approach, which may contain more complete attC sites than do shotgun reads—using SOAPdenovo (18) with k = 39 (see below for the selection of k-mer parameter), producing both contigs and the assembly graph (a de Bruijn graph) (26) (Fig. 2B); (ii) searching for attC sites in contigs using BLAST (with an identity threshold of 70% and coverage threshold of 50%) and tagging contigs with attC sites to be used as constraints to constrain the next step; (iii) extracting paths that start from one tagged contig and end at another tagged contig using a depth-first search algorithm and assembling the sequences for each path; the maximum length from one integron attC site to another attC site is set to 5,000 bp (Fig. 2C); (iv) checking the support of each assembled sequence by mapping the reads and read pairs onto the assembled sequences using BWA (17); we consider that a traverse between two contigs is valid if the flanking regions of the connection (of 1 bp at both sides; l is set to 15) are supported by at least one read or read pair, and an assembled sequence is considered to be supported only if all the traverses involved are supported by reads (Fig. 2D); (v) predicting the genes in each assembled sequence using FragGeneScan (30), with error model turned off; we require that the maximum gene number between any two integron attC sites be 3, considering that most integron cassettes contain 1 to 3 genes (6) (Fig. 2E).

**Validation of constrained assembly using simulation.** We simulated three metagenomic data sets by sampling reads at different coverage (10X, 20X, and 31X) from nine *Treponema* genomes (or genome drafts) using MetaSim (32) with the Illumina 80-bp error model with an error rate of ~1% provided by the authors (http://ab.inf.uni-tuebingen.de/software/metasim/errormodel-80bp.mconf). The species include *T. denticola* ATCC 35405 (NC_002967), *T. azonatorum* ATCC 35405 (NC_015577), *T. primitia* ZAS-2 (NC_015578), *Treponema pallidum* subsp. pallidum SS14 (NC_010741), *T. pallidum* subsp. pallidum strain Nichols (NC_000919), *T. succinifaciens* DSM 2489 (NC_015385), *T. denticola* strain F0402 (downloaded from http://www.broadinstitute.org/), *T. vincentii* (http://hmppdacc.org), and *T. phagedenis* (http://hmppdacc.org). We tested different k-mer parameters for the constrained assembly approach using these simulated data sets, and the results show that k = 39 resulted in the most integron genes for all the data sets (Fig. 3). The 31X data set contains 4,499,532 paired-end reads and 500,468 singleton reads. Seventy-three integron genes were identified from this data set by our constrained assembly approach: 37 genes from *T. denticola* ATCC 35405, 27 genes from *T. denticola* strain F0402, seven genes from *T. vincentii*, and two genes from *T. phagedenis*. We mapped these genes back to the genomes and confirmed that (i) all the genes were correctly assembled (error rate is 0%), and (ii) all the genes were mapped to the big integron or the degenerate, small integron region in the genomes (see Results). In addition, we did not find any genes in the *Treponema* species that do not harbor integrons. All suggest that our constrained approach is reliable even when reads from closely related species are present.

**Functional annotation of identified gene cassettes.** We downloaded all protein sequences from the eggNOG v2.0 database (23) and retrieved the sequences with COG annotation (40). MUSCLE (11) was used to generate a multiple alignment for each COG family, and the HMM builder from the HMMER3 package (10) was then applied to build an HMM for each COG. HMMER searches (by hmmscan from the HMMER3 package) were used to annotate the predicted integron gene cassettes, with an E value cutoff of 0.001. For a gene with COG hits, we recorded the best nonoverlapped results, so that if a gene encodes multiple domains with distinct functions, all the functions will be reported.

**Identification of potential source species of gene cassettes.** We used MEGAN (15) to identify the possible source species of the identified gene cassettes. We searched the genes against the NCBI NR database (as of September 2011) using BLASTP and applied the MEGAN software to

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**FIG 2** A diagram of the constrained assembly approach. (A) Paired-end and singleton reads from a metagenomic data set. (B) Assembly of all reads using SOAPdenovo, to generate contigs and a de Bruijn graph that connects the contigs. (C) Identification of contigs that consist of integron recombination repeats (shown as orange bars) and search for paths that start and end at a contig with repeats, using a depth-first search algorithm. At any intermediate node, the process will sort the coverage of all contigs connected by its outgoing edges and begin searching from the highest one. The starting and ending contig could be the same contig. (D) Validation of the assembled sequences (the paths) by read mapping and discarding of the paths that are not supported by reads (e.g., the middle sequence in the figure is discarded). (E) Identification of the integron repeats and their exact locations in the assembled sequences. Prediction of genes using FragGeneScan. Output sequences are between two repeats (attC sites) and consist of three or fewer genes. (F) Retrieval of the genes from sequences that pass all criteria.
The number of integron genes discovered from simulated metagenomic data sets using different k-mer settings. The x axis lists the k-mers, while the y axis shows the total number of genes assembled. We generated three data sets with different coverages (10×, 20×, and 31×) and applied our constrained assembly method to these data sets. Lines indicate the gene numbers found, and dashed lines are the numbers of genes that were identified solely at the contig level (i.e., genes on the contigs that are bounded between two integron recombination sites).

Analyze the similarity search results. Since the average length of the genes is 506 bp, we set the minimum score threshold to 100, as suggested by MEGAN’s authors for longer reads.

**HMP data sets.** We used the Human Microbiome Illumina WGS Reads (HMGWS) Build 1.0 and the whole-metagenome assemblies (PGAs) from the HMP consortium (http://www.hmpdacc.org/). There are 757 total metagenomic samples from 103 subjects (individuals). The reference genomes were also downloaded from this website.

**Availability of software and predicted integron genes.** We implemented the integron discovery pipeline in C++ and Perl. Software and predicted integron genes can be downloaded from our website http://omicsinformatics.indiana.edu/integron.

**RESULTS**

The *T. denticola* integron attC sites are unique to *Treponema* species. BLAST searches using the eight representative attC sequences against the NCBI nucleotide collection (NT) and the genome database (chromosomes) with default settings hit only *Treponema* genomes. Using an identity threshold of 70% and coverage threshold of 50%, 64 attC sites were found in the *T. denticola* ATCC 35405 genome, of which 45 are located within the chromosomal integron (positions 1817049 to 1874294) identified by reference 6. We also found two additional attC sites downstream of the integron region, suggesting that the integron may be even larger and contain more genes. The attC site located immediately downstream of the previously reported integron location is more degenerative (barely passes the coverage threshold), but the site further downstream is more complete, and we believe that these two attC sites are genuine. In addition, we found 7 attC sites outside the big integron region (for example, there is an attC site located between positions 300167 and 300227, which shares 98% sequential identity with the attC site within the integron array between positions 1870410 and 1870474). Furthermore, a degraded intI gene exists between positions 302289 and 302350, suggesting that a degraded, small integron may exist in this region of the genome. We also discovered integron sequences in *T. denticola* F0421 (sequence downloaded from http://www.broadinstitute.org/). While the integrase genes (intI) are very similar between these two strains (with 95% identity), the integron gene cassettes are quite different—only 10 integron genes are shared between these two strains (see Fig. S1 in the supplemental material).

Instances of the *T. denticola* attC sites were also found in the draft assemblies of two human microbiome reference genomes (as of July 2011): *T. vincentii* ATCC 35580 and *T. phagedenis* F0421. A total of 16 attC sites were found in five contigs of *T. vincentii* ATCC 35580, and 6 attC sites were found in three contigs of *T. phagedenis* F0421. We further checked the *T. vincentii* and *T. phagedenis* genomes for features indicative of integrons. In both genomes, there are gene cassettes flanked by attC sequences: we identified one gene in a *T. phagedenis* contig and 12 genes from three contigs of *T. vincentii*. One of the *T. vincentii* contigs exhibits a very clear integron structure, as shown in Fig. S2 in the supplemental material. None of the 12 genes identified in *T. vincentii* shared significant similarity with the integron genes of the *T. denticola* integron, suggesting that the gene cassettes of the two integron loci have undergone substantial changes since these two species diverged. We also searched the *T. vincentii* and *T. phagedenis* genomes using the *T. denticola* intI gene and detected a significant (sequence similarity = 86%) and (long(953-bp) intI gene on the *T. vincentii* contig ACYH1000073, which is shown in Fig. S2. Together with the recombination sites and the gene cassettes, this region contains all elements required for an integron.

**Integron-containing *Treponema* species are found in 80% of a normal human population, using searches for integron attC sites.** We identified integron attC sites in 300 of >700 HMP samples, using targeted assembly. The body sites that have identified integrons are summarized in Table 1. Most samples with integrons are orally related (including hard palate, supragingival plaque, saliva, tongue dorsum, subgingival plaque, throat, buccal mucosa, and attached/keratinized gingiva sites), whereas non-oral samples, including stool and vaginal samples, do not contain integron attC sites (repeats). This evidence suggests that a high proportion of oral samples contain the *Treponema* species implicated in dental diseases, implying that these pathogens are ubiquitous among people. The existence of *Treponema* species implicated in dental diseases in most normal human individuals (though of low abundance) is also supported by mapping the sequencing reads onto the available complete genomes (or drafts) of the three integron-containing *Treponema* species.

<table>
<thead>
<tr>
<th>Location</th>
<th>No. of samples with attC sites</th>
<th>Total no. of samples</th>
<th>% of samples with attC sites</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hard palate</td>
<td>1</td>
<td>1</td>
<td>100</td>
</tr>
<tr>
<td>Supragingival plaque</td>
<td>98</td>
<td>128</td>
<td>77</td>
</tr>
<tr>
<td>Saliva</td>
<td>5</td>
<td>5</td>
<td>100</td>
</tr>
<tr>
<td>Tongue dorsum</td>
<td>109</td>
<td>136</td>
<td>80</td>
</tr>
<tr>
<td>Vaginal introitus</td>
<td>0</td>
<td>5</td>
<td>0</td>
</tr>
<tr>
<td>Stool</td>
<td>0</td>
<td>150</td>
<td>0</td>
</tr>
<tr>
<td>Midvagina</td>
<td>0</td>
<td>2</td>
<td>0</td>
</tr>
<tr>
<td>Subgingival plaque</td>
<td>8</td>
<td>8</td>
<td>100</td>
</tr>
<tr>
<td>Throat</td>
<td>6</td>
<td>7</td>
<td>86</td>
</tr>
<tr>
<td>Posterior fornix</td>
<td>0</td>
<td>62</td>
<td>0</td>
</tr>
<tr>
<td>Anterior nares</td>
<td>2</td>
<td>94</td>
<td>2</td>
</tr>
<tr>
<td>Buccal mucosa</td>
<td>60</td>
<td>122</td>
<td>49</td>
</tr>
<tr>
<td>Right retroauricular crease</td>
<td>2</td>
<td>18</td>
<td>11</td>
</tr>
<tr>
<td>Left retroauricular crease</td>
<td>0</td>
<td>9</td>
<td>0</td>
</tr>
<tr>
<td>Palatine tonsils</td>
<td>6</td>
<td>6</td>
<td>100</td>
</tr>
<tr>
<td>Attached/keratinized gingiva</td>
<td>3</td>
<td>6</td>
<td>50</td>
</tr>
</tbody>
</table>
Many more integron gene cassettes can be identified in the HMP data sets using constrained assembly. Using the whole-metagenome assemblies, we were able to retrieve only 66 integron-associated genes (see above). Application of our constrained assembly approach (see Materials and Methods for details and a validation of the method using simulated data sets) to the HMP data sets led to the identification of 794 genes in 47 samples. After combining both predictions and keeping only unique genes for each sample, we derived a total of 826 unique genes (598 97% nonredundant). Table S1 in the supplemental material shows the comparison of the results by the constrained assembly approach results and the whole-metagenome assembly for individual HMP samples. The distribution of sample locations and the number of genes in each location are listed in Table S2. We identified genes in 24 supragingival plaque samples, 19 tongue dorsum samples, and 4 subgingival samples. The proportion of samples with gene cassettes identified using the constrained assembly approach is still low—compared with samples with identified attC sites (300)—due to the low abundance of the Treponema species in many samples (see Fig. S3). But we can still utilize the attC sites (taking advantage of the multiple copies of the attC sites) to identify T. denticola or related species in those samples, demonstrating the power of using unique repeats to trace rare species. We note that mapping shotgun sequences onto the known reference genomes (or drafts) of Treponema species can be used to identify the existence of these species in the HMP samples, but such a mapping cannot be effectively used to identify the integron gene cassettes due to the dynamic nature of the integron genes (e.g., the two T. denticola strains share only 10 cassette genes; see above).

Similarly, among the 300 samples with detected attC sites, the samples with gene cassettes assembled by constrained assembly were significantly larger than those with no identified genes (Welch’s t test, Z = 4.42, degrees of freedom = 68, P < 0.001). This can also explain why we did not find gene cassettes in samples from buccal mucosa—the buccal mucosa samples are significantly smaller than other oral data sets (Welch’s t test, Z = 25.28, degrees of freedom = 388, P << 0.001), partially caused by a large contamination of human DNAs in the buccal mucosa samples.

**Many cassettes are of unknown function.** We annotated the predicted cassette genes using similarity searches. Among the 826 genes, 501 cannot be assigned to a COG family (Table 2): ~60% are unassigned. Of the remaining genes, ~60% are assigned to COG categories R (general function prediction only) and S (function unknown). Combining these two categories, 85% of the 826 genes are of unknown function: the proportion is even higher than reported for other integrons (it was reported that 75% of the cassette pool associated with Vibrionales genomes corresponds to genes with undefined functions [4, 5]).

To analyze genes with identified functions, we clustered the genes within each location (at 97% identity) to see how many

**TABLE 2 Breakdown by body location of the samples that have identified Treponema chromosomal integron gene cassettes**

<table>
<thead>
<tr>
<th>Body location</th>
<th>No. of samples</th>
<th>No. of genes</th>
<th>No. of genes without COG hits</th>
</tr>
</thead>
<tbody>
<tr>
<td>Supragingival plaque</td>
<td>24</td>
<td>457</td>
<td>252</td>
</tr>
<tr>
<td>Tongue dorsum</td>
<td>19</td>
<td>283</td>
<td>203</td>
</tr>
<tr>
<td>Subgingival plaque</td>
<td>4</td>
<td>86</td>
<td>46</td>
</tr>
</tbody>
</table>
genes are unique to distinct locations. The functional category L (replication, recombination, and repair) is the majority among all functional categories (25%); genes associated with categories D (cell cycle control, cell division, and chromosome partitioning), K (transcription), N (cell motility), and T (signal transduction mechanisms) are also elevated among all functional categories, with 12%, 11%, 13%, and 13% of the genes with known functions, respectively (see Table S2 in the supplemental material). Integron genes with these functions have been reported previously: for example, category L and category T are among the most prevalent functions reported by reference 5. Genes in other categories, such as genes predicted to be part of the toxin/antitoxin system in category D, DNA-methyltransferase in category K, and methyl-accepting chemotaxis protein in category N, were also reported by reference 5. This again demonstrates that our results are consistent with the previous findings of gene functions encoded by chromosomal integrons.

We further compared the predicted genes found in the HMP data sets with the genes in the T. denticola chromosomal integron (located at positions 1817049 to 1874294 on NC_002967, as reported by reference 6) using BLAST with an E value cutoff of 0.001. We found that of the 826 genes, 192 (23%) hit to the genome’s integron genes. We also found that of the 70 integron genes identified in the T. denticola genome, 39 (6%) genes are assigned to T. vincentii, and 262 genes likely originate from other species: 117 (14%) genes from other spirochete species and 145 (17%) genes from nonspirochete species. Table S3 in the supplemental material lists the candidate donor species, and the annotations of the potential donor genes in these species. Here, we show two examples: the first example is 14 genes assigned to the order Clostridiales, which were first discovered in soil but also appear in human microbiomes (14, 35), and the other example is 25 genes assigned to Spirochaeta caldaria, a thermophilic bacterium (27).

Most integron genes are unique to samples and individuals. In order to characterize the cassette genes shared among different samples, we clustered genes from different samples using CD-HIT (19), with an identity cutoff of 70% at the amino acid level, and then mapped the clustered genes to samples. Figure 6 clearly shows that gene sharing among samples is minimal. Most of the genes uniquely belong to only one sample—only 84 genes are shared between exactly two samples and 63 genes are shared among three or more samples. This finding is consistent with the findings from reference 16 that integron genes from 12 Vibrio isolates share only a very small number (<10%) of genes. The HMP cohort contains individuals who were sampled at multiple body sites and visits, enabling us to compare the sharing of the integron cassette genes within and across individuals. We calculated the proportion of shared genes between any two samples and found that samples from the same individual tend to share more genes than do samples from different individuals: the average proportion of genes shared between samples from the same individual is 13%, and the average proportion of genes shared between samples from different individuals is slightly lower, 8%. Note again, it was reported previously that Vibrio isolates share <10% of their integron genes (16). Our results indicate that even within an individual, there is strong population subdivision between Treponema species collected at different sites.

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**FIG 5** Taxonomic assignments of the integron genes by MEGAN. The numbers following clade names are the numbers of genes assigned to that taxonomic rank, not including the genes assigned to the taxa below that rank (for example, there are 63 genes assigned to T. denticola species, 49 genes assigned to strain ATCC 35405, and 40 genes assigned to strain F0402; in total, 138 genes can be assigned to the T. denticola species).
The functions of the shared genes also vary, and the majority of them are still of unknown function: for the 84 genes shared between two samples, 56 genes cannot be assigned to any COG function, and 19 are assigned to unknown function (category R or S). Similarly, for the 63 genes shared by three or more samples, 30 genes do not hit to any COG function and 17 genes hit to unknown functions. (See Table S4 in the supplemental material for detailed numbers of hits to COG functions.) Overall, the percentage of shared genes with an unknown function is 83%. This number is similar to the proportion for all 826 genes. Furthermore, the number of genes in category L (replication, recombination, and repair) is again the highest among all categories with known functions. These numbers hint that the genes shared among two or more samples are sampled from all integron genes, without any preference for genes of certain functions.

**DISCUSSION**

To assemble integron gene cassettes, we designed a novel method to trace the de Bruijn assembly graph and then extract sequences bounded by contigs that contain \textit{attC} sites. Assembly approaches based on de Bruijn graphs typically report the sequences of the edges (i.e., contigs) while discarding the connections between contigs embedded in the graph—the ambiguous connections between contigs may be difficult to resolve if no further information can be applied (28). Our novel constrained assembly approach to integron gene cassettes enables us to traverse between the contigs in the de Bruijn graph by applying further information learned from the integron structures. The effect is enormous, as we obtained 826 genes \textit{de novo} using this approach, compared to only 66 genes in the whole-metagenome assembly contigs.

Our integron gene discovery pipeline (see Materials and Methods) includes two validation steps (step 4 and step 5): only genes included in the sequences that are supported by read mapping (step 4) and in sequences that contain 1 to 3 genes (step 5) will be reported as candidate integron genes. For the HMP data sets, only 22% of sequences passed the first validation process and 56% of genes passed the second. We did not observe any misassembled integron genes when we applied the pipeline to the simulated data sets. We cannot completely exclude the possibility of having misassemblies in the real HMP data sets, considering that the prediction of the integron genes may be affected by reads from unknown species. Also, our method may miss some integron genes due to the heterogeneity of \textit{attC} sites of the \textit{Treponema} species in the real samples.

Our targeted assembly and constrained assembly approaches can in principle be applied to any metagenome con-
taining an integron system. Given the $attC$ sites, we are able to detect species with the corresponding integrons and generate integron gene cassettes. For example, the coral-mucus-associated *Vibrio* integrons (16) can be used to detect this coral pathogen in ocean samples, such as the Sargasso Sea metagenomic samples (41). By analyzing integron genes, we can help to understand how this species evolves and coexists with coral. We can also analyze genes from different sites (or depths) of the ocean and understand how bacteria in these sites interact with the outer environment. Even if species with integrons are of low abundance, we can still detect their existence in metagenomic samples, as in the case of *T. denticola*.

Note that our targeted assembly (used in this work to characterize the integron $attC$ sites) was developed to characterize CRISPR arrays in metagenomic samples (31). CRISPR/Cas systems are a widespread class of adaptive immunity systems that bacteria and archaea mobilize against foreign nucleic acids; the CRISPR arrays contain repeats, and short spacers that are likely derived from viral genomes or plasmids. Because the spacers in CRISPR arrays are significantly shorter than Illumina reads, we could easily assemble CRISPR arrays using targeted assembly alone, by first collecting reads containing repeats and then assembling the reads using optimized parameters. In contrast, integron spacers (cassettes) contain 1 to 3 genes between the $attC$ sites, so it is hard to assemble the gene cassettes using targeted assembly alone. The constrained assembly approach was developed to overcome this limitation and allows the assembly and characterization of integron gene cassettes. Both applications (the identification of the CRISPR arrays using the targeted assembly approach and the identification of integron gene cassettes) demonstrate the importance of directed computational approaches for studies of important functional elements—which are poorly analyzed using generalized computational approaches (such as whole-metagenome assembly)—and show that they are essential for the analysis of metagenomic sequences.

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REFERENCES


