Comparative Analysis of AbaR-Type Genomic Islands Reveals Distinct Patterns of Genetic Features in Elements with Different Backbones

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ABSTRACT AbaR-type genomic islands (AbaRs) are prevalent and associated with multiple antimicrobial resistance in Acinetobacter baumannii. AbaRs feature varied structural configurations involving different but closely related backbones with acquisition of diverse mobile genetic elements (MGEs) and antimicrobial resistance genes. This study aimed to understand the structural modulation patterns of AbaRs. A total of 442 intact AbaRs, including nonresistance but closely related islands, were mapped to backbones Tn6019, Tn6022, Tn6172/Tn6173, and AbGRI1-0 followed by alien sequence characterization. Genetic configurations were then examined and compared. The AbaRs fall into 53 genetic configurations, among which 26 were novel, including one Tn6019-type, nine Tn6022-type, three Tn6172/Tn6173-type, nine AbGRI1-type, and four new transposons that could not be mapped to the known backbones. The newly identified genetic configurations involved insertions of novel MGEs like ISAcp2, ISAba42, ISAba17, and ISAba10, novel structural modulations driven by known MGEs such as ISCR2, Tn2006, and even another AbaR, and different backbone deletions. Recombination events in AbGRI1-type elements were also examined by identifying hybrid sequences from different backbones. Moreover, we found that the content and context features of AbaRs including the profiles of the MGEs driving the plasticity of these elements and the consequently acquired antimicrobial resistance genes, insertion sites, and clonal distribution displayed backbone-specific patterns. This study provides a comprehensive view of the genetic features of AbaRs.

IMPORTANCE AbaR-type genomic islands (AbaRs) are well-known elements that can cause antimicrobial resistance in Acinetobacter baumannii. These elements contain diverse and complex genetic configurations involving different but related backbones with acquisition of diverse mobile genetic elements and antimicrobial resistance genes. Understanding their structural diversity is far from complete. In this study, we performed a large-scale comparative analysis of AbaRs, including nonresistance but closely related islands. Our findings offered a comprehensive and interesting view of their genetic features, which allowed us to correlate the structural modulation signatures, antimicrobial resistance patterns, insertion loci, as well as host clonal distribution of these elements to backbone types. This study provides insights into the evolution of these elements, explains the association between their antimicrobial resistance gene profiles and clonal distribution, and could facilitate establishment of a more proper nomenclature than the term “AbaR” that has been variously used.
Multidrug-resistant *Acinetobacter baumannii* remains a major threat of nosocomial infection. AbaR-type genomic islands (AbaRs) are a class of important mobile genetic elements (MGEs) prevalent in *A. baumannii* and have been reported to carry multiple antimicrobial resistance genes (1–4).

AbaRs exhibit variable genetic structural features involving certain different but closely related backbones and diverse acquired transposons, insertion sequences (IS), and antimicrobial resistance genes based on these genetic elements (5–8). Tn6019, Tn6022, and Tn6172/Tn6173 have been recognized as backbone elements (5, 8–10). Tn6173 is a hypothetical transposon from which Tn6172 is derived (8). These backbones share a 2.89-kb left-end conserved sequence (CS) and a 1.87-kb right-end CS spanning transposition genes (4). AbaRs commonly disrupt the chromosomal *comM* gene. Elements with different backbones are usually found in different epidemic clones. The well-documented AbaR3-type elements are seemingly confined to global clone 1 (GC1) (3). They contain a Tn6019 backbone and are invariably associated with Tn6018 or its compound elements containing plastic multiple antimicrobial resistance regions (MARRs) (3). The Tn6022-derived elements mostly are found in GC2 and sometimes bear the *blaOXA-23* gene as seen in AbaR4 (9). A class of complex elements called AbGRI1-type islands have recently been identified in GC2, which are proposed to originate from a plasmid-borne ancestral form (AbGRI1-0) consisting of a Tn6022, a Tn6172, and a plasmid-borne fragment between the two transposons (termed “linker” here) as a transposable unit (8, 12). Our recent large-scale identification of AbaRs in *A. baumannii* genomes have also uncovered that they have diverse insertion sites and clonal lineage-specific antimicrobial resistance gene profiles (4). Although efforts have been made to study the evolution of AbaRs (6, 13–15), understanding of their structural diversity is far from complete.

The term “AbaR” was initially coined for *A. baumannii* resistance islands (1). It has also been used for highly homologous islands without resistance genes that were discovered later. All these elements displayed wide diversity, but they could be genetically mapped to certain backbones as mentioned above, regardless of whether they carried resistance genes. There is still a lack of consensus on the nomenclature of these elements. For simplicity, here we collectively called these islands “AbaRs” (4). To avoid confusion, it should be noted that the term was used only to refer to these genetically related islands and was not limited to islands with resistance genes.

In this study, exhaustive profiling and comparative analysis of the genetic features of 422 intact AbaRs with complete genetic information were performed, which allows us to correlate their backbone types with structural modulation signatures, antimicrobial resistance patterns, insertion loci, as well as clonal distribution.

RESULTS

Twenty-six novel AbaR genetic configurations were identified. A total of 442 AbaRs were selected (see Data Set S1 in the supplemental material), among which 65 nonredundant elements were identified. They were further mapped to backbones Tn6019 (5), Tn6022 (9), Tn6172 (10), Tn6173 (8), and AbGRI1-0 (8), followed by comparative analysis. AbaRs were characterized into 53 genetic configurations (see Data Set S1 and Table S1 in the supplemental material). Three configurations were present as intact backbones, including Tn6019, Tn6022, and Tn6172, 46 were variants associated with the known backbones, while four (designated Tn6661 to Tn6664) were new transposons that could not be mapped to any of the known backbones. Of note, 26 configurations were novel ones, including one Tn6019-derived, nine Tn6022-derived, three Tn6172/Tn6173-derived, nine AbGRI1-type, and four new transposons.

Comparative analysis of Tn6019-derived elements. Five configurations were found to be based on Tn6019 (Fig. 1). Four of the configurations were associated with Tn6018 or its compound elements that disrupted the *uspA* gene and have been widely...
seen and well documented (3). We found a novel configuration that had an IS\textsc{Aba}1 disrupting orf4, but there was only one element bearing this configuration. Note that AbaRs containing the Tn\textsc{6018} compound elements were characterized as one configuration (type) here for simplicity despite the variable nature of the internal MARRs (3), unless there were subsequent deletions on the backbones.

**Comparative analysis of Tn\textsc{6022}-derived elements.** Thirteen configurations were characterized as Tn\textsc{6022} derived (Fig. 2), with variations involving insertions of diverse MGEs and backbone deletions. The IS\textsc{Aba}11 and Tn\textsc{2006} (IS\textsc{Aba}1-bounded composite transposon) were frequently seen and site specifically disrupted \textit{tniC} and \textit{sup}, respectively. Insertions of IS\textsc{Acsp}2, IS\textsc{Aba}42, and IS\textsc{Aba}1 were newly found. IS\textsc{Acsp}2 disrupted \textit{tniB}, the IS\textsc{Aba}42 inserted into the intergenic region between \textit{tniE} and \textit{orf}, while IS\textsc{Aba}1 was seen at different sites. Four novel deletion forms of the backbone were found in addition to Tn\textsc{6022}\textsc{Δ1} (2). The Tn\textsc{6022}\textsc{Δ3} had an extra 1.49-kb deletion based on Tn\textsc{6022}\textsc{Δ1}, while in Tn\textsc{6022}\textsc{Δ2}, Tn\textsc{6022}\textsc{Δ4}, and Tn\textsc{6021}\textsc{Δ5}, different internal fragments (0.6 to 4.6kb) were missing. Note that the deletion in Tn\textsc{6022}\textsc{Δ2} likely involved 6-bp perfect duplicated target sequences (5\textsc{′}-\textit{AAATGC}-3\textsc{′}), suggesting the missing fragment might be a novel IS.

**Comparative analysis of AbGRI1-type elements.** Twenty-two complex configurations containing more than one backbone were all characterized as AbGRI1-type islands (Fig. 4 and 5). They displayed complex chimeric mosaicism. They had expanded variation profiles other than those of the solitary Tn\textsc{6022}- and Tn\textsc{6172}-derived elements (Table S2). In addition, IS\textsc{Aba}17 and IS\textsc{Aba}10 were found on the Tn\textsc{6022}-derived
portions. Inversions of Tn2006 were also observed. As for the Tn6173-derived part, there could be an insertion of a second Tn6022, Tn6022-derived, or even an AbGRI1-type element, splitting the 3’ end of the tet(B) gene that is located on the ΔISCR2-ΔTn10 segment, consistent with our previous observation that tet(B) was a high-frequency

| Name          | Genetic feature                                                                 |
|---------------|---------------------------------------------------------------------------------|
| Tn6022        | backbone                                                                         |
| Tn6021        | Tn6022ΔtniC::IASba11                                                              |
| AbaR4         | Tn6022Δsup::Tn2006                                                                |
| NA            | Tn6022ΔtniB::IAScsp2                                                               |
| NA            | Tn6022(tniE-orf4)::IASba42                                                        |
| Tn6022Δ1      | Tn6022 with ~2.85-kb internal deletion                                           |
| Tn6022Δ1      | (named by this study) Tn6022Δ1ΔtniC::IASba11                                      |
| NA            | Tn6022Δ1ΔtniB::IASba1                                                           |
| NA            | Tn6022Δ1[ΔtniC::IASba11,(sup-orf4)::IASba1]                                      |
| RIA2R1//AbaR4Δ1| (re-named by this study) Tn6022Δ1Δsup::Tn2006                                     |
| Tn6022Δ3      | Tn6022Δ1 with ~1.49-kb internal deletion                                         |
| Tn6022Δ2      | Tn6022 with 612-bp internal deletion                                           |
| Tn6022Δ4      | Tn6022 with 1.98-kb internal deletion                                            |
| Tn6021Δ5      | Tn6021 with ~4.58-kb deletion                                                   |

**FIG 2** Comparison of Tn6022-derived genetic configurations. The Tn6022 backbone is shown at the top of the figure. The backbone segments in the variants are placed in aligned positions. The names of the reported elements bearing the corresponding genetic configurations are given. NA, not available. The names given by this study are also shown. Novel genetic configurations are shown in red. Dotted lines are used to connect sequences. The schematic representations are drawn to scale.
We also found that the ΔISPa14-Tn1213-strA region on the Tn6173 part of AbaR4d (7) was variable, associated with the acquisition of aph(3')-VIb. Moreover, a considerable fraction of AbGR11-type configurations (Fig. 5) did not possess a typical Tn6022-linker-Tn6172 structure, since the CSS from different backbones provide targets for homologous recombination, as illustrated by AbGR11-5 (16). AbGR11-5 featured a hybrid region made up of Tn6022 and Tn6173 segments. We further uncovered similar phenomena on AbaR26(BJAB0868) (17), AbaR25(BJAB07104) (17), and AbaR22 (18) (Fig. 5; see also Fig. S1 in the supplemental material). However, the precise boundaries of segments from different backbones were not specific and could hardly be defined. A hybrid region was not found on other atypical elements like AbaR4a/AbaGR11-1, AbaR4b, and AbaR4c (7).
Structures of Tn6661, Tn6662, Tn6663, and Tn6664. Tn6661 (12.2 kb), Tn6662 (13.1 kb), Tn6663 (20.0 kb), and Tn6664 (23.2 kb) were distinct elements (Fig. 6) showing 71% to 92% nucleotide identities to the known backbones (Fig. S2) and each other (Fig. S3) at the syntenic regions, covering the CSs at both ends. Like other backbone elements, they carried distinct genes in the middle regions. However, no antimicrobial resistance gene was identified. The Tn6661 transposon harbors predicted \textit{crcB} and \textit{ppa} genes that might function as a fluoride transporter and an inorganic pyrophosphatase, respectively. Tn6662 harbors genes that might participate in biosynthesis, as they were predicted to encode a PhzF family phenazine biosynthesis protein, a transporter, an oxidoreductase, and a transcriptional regulator. Tn6663 carries a \textit{cop} gene cluster likely involving in copper resistance. Tn6664 carries a set of genes that might be linked to...
FIG 5 Genetic configurations of the AbGRI1-type variants that might have gone through recombination. The proposed ancestral form (AbGRI1-0) of AbGRI1-type elements is shown at the top of the figure. An example of a proposed recombination process is shown (see AbGRI1-3 and AbGRI1-5) (16). The syntenic regions with near 100% identity between AbGRI1-3 and AbGRI1-5 are connected by gray. The Tn6022 or ΔTn6022 part is shown in orange; the Tn6172 part or its partial segments are shown in blue, while the linker region is shown in green. Where possible, the backbone segments in the variants are placed (Continued on next page)
certain biochemical process, as they encoded putative enzymes such as disulfide reductase, dehydrogenase, oxidase, decarboxylase, and protein-disulfide reductase. Apart from an IS17-like element found on Tn6662, no other MGE was found on these elements, suggesting that they could serve as backbone structures. Interestingly, we found a Tn6664-type variant in Acinetobacter haemolyticus strains TJS01 (GenBank accession number CP018871; element coordinates, 3136595 to 3162983) (19). It had an ISAba8-like IS and an ISscp3-like IS based on Tn6664. However, other variants based on these elements are yet to be revealed.

The content and context characteristics of AbaRs were associated with backbones. Finally, heatmaps profiling the acquired MGE pools, antimicrobial resistance gene profiles, insertion sites, and clonal distribution for the nonredundant AbaRs were generated (Fig. 7) and displayed apparent backbone-specific patterns. As mentioned above, the Tn6019-, Tn6022-, Tn6172/Tn6173-, and AbGRI1-type variants were associated with different sets of acquired MGEs. Moreover, they carried distinct spectrums of antimicrobial resistance genes that were specific to the backbones. They also showed differences in insertion sites and clonal distributions. Consistent with previous reports (3, 16), the Tn6019- and AbGRI1-type elements were mostly associated with the conventional comM insertion site, with very few exceptions in other loci, and almost confined to specific clonal lineages, GC1 and GC2, respectively. However, the Tn6022-type AbaRs had more diverse insertion sites and wider clonal distribution (Fig. 7 and Fig. S4) than the other types. The Tn6172/Tn6173-type AbaRs were associated with a plasmid-borne insertion site instead of the comM. The Tn6661, Tn6662, Tn6663, and Tn6664 transposons were found in minor clones, and their features need further investigation, as there might be sample bias in the current genome database.

**DISCUSSION**

AbaRs are prevalent in *A. baumannii*. We had previously performed large-scale identification of AbaRs in *A. baumannii* genomes and found that their antimicrobial resistance gene profiles were specific to clonal lineage (4). However, the exact genetic configurations involving backbone types and variation patterns of the identified elements remained unknown. Here, we conducted further comparative genomic analysis of AbaRs. We uncovered novel genetic configurations of AbaRs and found that the content and context characteristics of AbaRs were specific to backbones.

The newly identified genetic configurations mainly involved insertions of novel MGEs or novel structural modulations driven by known MGEs. We found a rare

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**FIG 5 Legend (Continued)**

in aligned positions. The names of the reported elements bearing the corresponding genetic configurations are given. NA, not available. Novel genetic configurations are shown in red. Dotted lines are used to connect sequences. Red blocks highlight the regions that might have gone through recombination, since they contained hybrid sequences from different backbones (for details, see Fig. S1 in the supplemental material). The schematic representations are drawn to scale.
Tn6019-derived configuration that carried an ISAba1. ISAba1 is an active IS in *A. baumannii* and has been widely found in this species. As for AbaRs, the ISAba1 has been seen on AbaR4, Tn6172, and some AbGRI1-type elements as part of Tn6006 or alone (5, 7, 20) but has not yet been reported as occurring on Tn6019. Tn2006 is closely associated with the variations based on Tn6022, as it has been frequently seen on Tn6022-type elements or the Tn6022-derived parts of AbGRI1-type elements (7, 13, 21, 22). Meanwhile, diverse backbone deletions were found on Tn6022, but the mechanisms of the variations remain unknown. ISCR2 might play a role in modulating the plasticity of Tn6172 and Tn6173, since the ΔISCR2-ΔTn10 fragment or a larger fragment containing that structure was often found at the ISCR2 locus (8). Moreover, in AbGRI1-type elements, the tet(B) gene on the ΔISCR2-ΔTn10 fragment provided a hot spot for insertion of another AbaR, resulting in complex chimeric structures. This type of configuration was found on reported elements like Rl$_{MDT_T}$/AbaR1 (14) and AbaR26(BJAB0868) (17), but it had not been previously elucidated. It seems the multiple homologous backbone CSS in the AbGRI1-type elements have actively triggered complex recombination events, generating diverse atypical AbGRI1-type elements containing hybrid backbone sequences. Although the genetic events behind this process remain unclear, it is undoubted that this nature could dramatically complicate the evolution of AbGRI1-type elements.
A comprehensive profiling of the genetic features of AbaRs uncovered backbone-specific patterns. The acquired MGE pools of AbaRs remarkably differed by backbone types, which directly resulted in distinct antimicrobial resistance gene profiles, since the acquired resistance determinants were almost associated with the acquired MGEs. For example, the multiple antimicrobial resistance of Tn6019-type AbaRs was attributed to the Tn6018 compound elements (3), the only resistance gene that appeared on Tn6022-type AbaRs, bla\textsubscript{OXA-23}, was embedded in Tn2006 (23), and the tet(B), bla\textsubscript{PER-1}, or other resistance genes on Tn6172/Tn6173 were cargo of ΔTn10, Tn1213, or ISCR2-associated regions, respectively (7, 8). Notably, on the CS regions, AbaRs still had backbone-specific structural variations. The variation might be due to the fact that the CSs of different backbones had nucleotide variations, some of which might result in favorable targeting sequences for certain MGEs, or to the possibility that the AbaRs might have experienced different MGE exposure in different clones. Indeed, AbaRs of most backbone types are associated with specific clonal lineages. We have previously observed the association between antimicrobial resistance gene profiles and clonal distribution (4); here, we were able to extend the correlation with backbone types. We have also reported that AbaRs have diverse insertion sites other than \textit{comM} (4). Here, we further demonstrated that this feature was mainly found in Tn6022-type AbaRs. In addition, it is likely that plasmids might promote the interclonal exchange of AbaRs (24), since the Tn6022-type and Tn6172/Tn6173-type AbaRs spanning multiple clonal lineages (Fig. 7; see also Fig. S4 in the supplemental material) both had plasmid-borne insertion sites. However, the AbGRI1-type islands did not share this feature and were mostly found in GC2. These results indicated that AbaRs with different backbones might have evolved separately.

The term “AbaR” initially referred to resistance islands in \textit{A. baumannii} (1) but has also been used for the closely related \textit{comM}-associated elements described above, regardless of whether they carry a resistance gene. It also should be noted that these elements are not confined to just the \textit{comM} locus. The term “AbaR” has certain limitations, since these elements display wide diversity. It would be better if the nomenclature were based on the transposition module and/or the transposition process of these elements. Given the backbone-specific nature of these elements revealed by this study, we propose that the elements with different backbones be treated separately, and that using backbone-based nomenclature or specified Tn (transposon) numbers could be more specific and informative than the generic term “AbaR” to reflect the genetic features of these elements. We envision that this study could help lead to a solution on the nomenclature of these elements.

Overall, this study uncovered genetic features of “AbaRs,” including the profiles of the MGEs driving the plasticity of these elements and the consequently acquired antimicrobial resistance genes, clonal distribution, as well as insertion sites displaying strong associations with their backbones.

**MATERIALS AND METHODS**

**Genomic data of AbaRs.** Information on 468 intact AbaRs from our recent identification of \textit{A. baumannii} genomes (4), literature, and unpublished entries in the GenBank database were collected, and 442 were included for analysis (see Data Set S1 in the supplemental material). Among the excluded AbaRs, 21 contained large sequencing gaps, and five turned out to be portions of AbGRI1-type elements with incomplete sequences rather than individual elements. Specifically, the last five were initially identified on contigs as intact elements (Tn6022) (4), but further analysis here showed that their upstream and downstream sequences could be mapped to the Tn6172 backbone, which meant that the identified elements were actually parts of AbGRI1-type elements. There was not enough sequence information on the corresponding contigs to recognize the entire elements with the previously described algorithm (4). Among the included AbaRs, 36 have been previously characterized, 402 were recently predicted, and 4 were from unpublished GenBank entries (Data Set S1). The latter two kinds have not been characterized before.

**Comparative analysis and genetic configuration profiling of AbaRs.** AbaRs were mapped to backbones Tn6019, Tn6022, Tn6172, Tn6173, and AbGRI1 via BLAST with a cutoff of >95% overall nucleotide identities (or >90% for elements with the Tn6022ΔΩ2 configuration). Alien sequences were characterized as previously described (11). Genetic configurations were defined manually. Complex elements \((n = 253)\) containing multiple backbones were also aligned with the elements \((n = 189)\) containing one backbone for comparison. A nonredundant set of the AbaRs was generated.
with a cutoff of >99% coverage and >98% identity (or >95% for the Tn6022Δ2 configuration) for analysis.

Multiple-sequence alignment. Multiple-sequence alignment was performed with MUSCLE (25) to identify hybrid regions from different backbones.

Heatmaps. Heatmaps were generated with the ComplexHeatmap package (26) in R.

Accession numbers. The accession numbers of all sequences used in this study are listed in Data Set S1 in the supplemental material along with precise coordinates where applicable.

SUPPLEMENTAL MATERIAL

Supplemental material is available only online.

FIG S1, TIF file, 1.1 MB.

FIG S2, TIF file, 0.9 MB.

FIG S3, TIF file, 0.5 MB.

FIG S4, TIF file, 1.1 MB.

TABLE S1, DOCX file, 0.1 MB.

TABLE S2, DOCX file, 0.02 MB.

DATA SET S1, XLS file, 0.2 MB.

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We declare that we have no conflicts of interest.

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