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Structure and Evolution of Chlorate Reduction Composite Transposons

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ABSTRACT The genes for chlorate reduction in six bacterial strains were analyzed in order to gain insight into the metabolism. A newly isolated chlorate-reducing bacterium (Shewanella algae ACDC) and three previously isolated strains (Ideonella dechloratans, Pseudomonas sp. strain PK, and Dechloromarinus chlorophilus NSS) were genome sequenced and compared to published sequences (Alicyclobacillus denitrificans BC plasmid pALIDE01 and Pseudomonas chloritidismutans AW-1). De novo assembly of genomes failed to join regions adjacent to genes involved in chlorate reduction, suggesting the presence of repeat regions. Using a bioinformatics approach and finishing PCRs to connect fragmented contigs, we discovered that chlorate reduction genes are flanked by insertion sequences, forming composite transposons in all four newly sequenced strains. These insertion sequences delineate regions with the potential to move horizontally and define a set of genes that may be important for chlorate reduction. In addition to core metabolic components, we have highlighted several such genes through comparative analysis and visualization. Phylogenetic analysis places chlorate reductase within a functionally diverse clade of type II dimethyl sulfoxide (DMSO) reductases, part of a larger family of enzymes with reactivity toward chlorate. Nucleotide-level forensics of regions surrounding chlorite dismutase (cld), as well as its phylogenetic clustering in a betaproteobacterial Clr clade, indicate that cld has been mobilized at least once from a perchlorate reducer to build chlorate respiration.

IMPORTANCE Genome sequencing has identified, for the first time, chlorate reduction composite transposons. These transposons are constructed with flanking insertion sequences that differ in type and orientation between organisms, indicating that this mobile element has formed multiple times and is important for dissemination. Apart from core metabolic enzymes, very little is known about the genetic factors involved in chlorate reduction. Comparative analysis has identified several genes that may also be important, but the relative absence of accessory genes suggests that this mobile metabolism relies on host systems for electron transport, regulation, and cofactor synthesis. Phylogenetic analysis of Cld and ClrA provides support for the hypothesis that chlorate reduction was built multiple times from type II dimethyl sulfoxide (DMSO) reductases and cld. In at least one case, cld has been coopted from a perchlorate reduction island for this purpose. This work is a significant step toward understanding the genetics and evolution of chlorate reduction.

Perchlorate (ClO4−) and chlorate (ClO3−) have natural and anthropogenic sources. While recent evidence suggests that these compounds are formed in the atmosphere (1, 2), contamination of drinking water is often a result of human activity. Chlorate has been used as an herbicide and defoliant and as a bleaching agent in the paper industry; perchlorate is a solid oxidant found in flares, explosives, and propellants (3). Bacterial remediation of contaminated water is a viable treatment option, which has spurred both applied (4) and basic (5) science research. Perchlorate and chlorate are respired by dissimilatory perchlorate-reducing bacteria (PRB) and chlorate-reducing bacteria (CRB), respectively, almost all of which are Proteobacteria (6), with a few exceptions (7, 8) (see Fig. S1 in the supplemental material). While all PRB isolated are also chlorate reducers, the reverse is not true. The distinction is at least partly a result of the specificity of the terminal reductase; the perchlorate reductase (PcrAB) can reduce perchlorate and chlorate (9), while the chlorate reductase (ClrABC) can reduce only the latter (10, 11). Chlorite is an obligate intermediate in both pathways and is detoxified by the chlorite dismutase (Cld), which produces chloride and molecular oxygen that is respired. The chlorate reductases of Ideonella dechloratans, Pseudomonas chloritidismutans AW-1, and Pseudomonas sp. strain PDA have been purified as soluble heterotrimers (αβγ) (10–12). ClrABC in I. dechloratans and that in PDA are probably periplasmic, and while fractionation experiments support a cytoplasmic ClrABC in AW-1, a twin-arginine signal motif is predicted (13), suggesting periplasmic localization. By comparison to structurally characterized enzymes EbdABC (14) and NarGHI (15), the α subunit is predicted to contain a bis(molybdopterin guanine dinucleotide)-molybdenum cofactor and a [4Fe-4S] cluster coordinated by one histidine and three cysteines (10). The β subunit is predicted to contain four Fe-S clusters that form an
electron transfer pathway between a cytochrome $b$ in the $\gamma$ subunit (16) and the Fe-S cluster in the $\alpha$ subunit. The $\delta$ subunit is homologous to NarJ and most likely participates in proper insertion of the molybdenum cofactor but is not part of the active enzyme (17, 18).

To date, three CRB (Ideonella dechloratans, P. chloritidismutans AW-1, and Alicyclobacillus denitrificans BC) have had their genes for chlorate reduction sequenced. As part of our continuing effort to understand the genomics of chloroxanion respiration by bacteria, genome sequences were completed for four CRB: Ideonella dechloratans, Pseudomonas sp. strain PK, Dechloromonas chlorophilus NSS, and the newly isolated Shewanella algae ACDC. However, after de novo assembly, the genes for chlorate reduction were found on small contigs, with no information about neighboring regions. Short reads from next-generation sequencing (NGS) technologies often do not unambiguously connect regions surrounding repeats, and as a result, assemblers produce many contigs instead of contiguous finished genomes. A fragmented genome may not be a research impediment if the genes of interest are on a large contig. This was the case in our recent comparative analysis of genes for perchlorate reduction, in which conserved synteny and evidence of horizontal gene transfer led to the identification of a perchlorate reduction genomic island (PRI) that contained metabolic, regulatory, and electron transport chain components (19).

Using a bioinformatics approach, contigs containing genes for chlorate reduction (clrABDC and cld) were extended, connected to neighbors, and confirmed with finishing PCRs. In all four newly sequenced chlorate-reducing bacteria, these genes are flanked by insertion sequences, forming composite transposons. Insertion sequences are small, autonomous, mobile genetic elements that encode a transposase and accessory genes enclosed by terminal repeat sequences (20). Insertion sequences are widely recognized as drivers of bacterial evolution; they are often present in large numbers and serve as sites for homologous recombination and therefore chromosomal rearrangements, plasmid integration, and deletions (21). As composite transposons, they construct and disseminate novel metabolic pathways, including those involved in the catabolic degradation of substituted aromatics and xenobiotics (22, 23). The identification of chlorate reduction composite transposons is an important step toward understanding the formation and horizontal transfer of this metabolism.

RESULTS

After de novo assembly, genome sequences of PK, NSS, and ACDC contained cld on a contig with a gene cluster encoding a dimethyl sulfoxide (DMSO) reductase family type II enzyme. This was inferred to be the chlorate reductase based on its proximity to cld and phylogenetic clustering with other putative chlorate reductases (10, 11). Since no genetic system is available, the physiological function of the chlorate reductase was obscured by fragmented assembly of NGS reads, which produced clrABDC and cld on contigs of less than 16 kb. To resolve neighboring genes, assemblies of I. dechloratans, PK, ACDC, and NSS were improved by mapping reads to the ends of contigs, extending them computationally (24), and predicting links to other contigs (see Materials and Methods).

Shewanella algae ACDC and Dechloromonas chlorophilus NSS. ACDC was isolated from the same marine sediment enrichment from which NSS was previously isolated (6). Details of the physiology of ACDC will be published elsewhere. After improved assembly and annotation (Fig. 1A), two ISPpu12 insertion sequences flanked clrABDC and cld (Fig. 1A). Five copies of this composite transposon were found in ACDC: two chromosomal insertions and one on a multicopy (~3) plasmid (Fig. 1D). The insertion sequences flanking the plasmid-borne copy are isoforms, designated ISPpu12a and ISPpu12b, and contain 68 single nucleotide polymorphisms in the transposase gene, as well as a small deletion of 12 bp in czd (Fig. 1B). The chromosomal composite transposons contain matching flanking copies of ISPpu12b. The reason for the discrepancy between plasmid and chromosomal insertion sequence isoforms is unknown but was the cause for negligible assembly of this region (Fig. 1C and D). Imperfect 24-bp inverted repeats are located at the boundaries of both ISPpu12 isoforms and are identical to those reported for ISPpu12 in the Pseudomonas putida plasmid pWW0 (Fig. 1E).

Insertion of the composite transposon into the ACDC chromosome occurred in the open reading frames of nrfA (ammonium-forming nitrite reductase) and barA (hybrid sensor kinase). This finding is supported by conserved synteny surrounding nrfA and barA in closely related species, by the continuous coverage of mapped reads over the insertions, and by PCRs connecting nrfA and barA to the interior of the composite transposon (Fig. 1C). Presumably due to the insertion in nrfA, ACDC reduces nitrate to nitrite but does not reduce nitrite to ammonium (I. C. Clark, unpublished data). Duplications of 8 bp found surrounding the composite transposon insertions in nrfA, barA, and the plasmid (Fig. 1E) have been previously observed in other ISPpu12 elements following transposition (25).

Assembly of the NSS genome produced contigs that matched ACDC’s plasmid with 99.9% nucleotide identity (Fig. 1D). The plasmid contains tra and trb gene clusters putatively involved in self-transmissibility. Unlike in ACDC, coverage of the composite transposon in NSS is consistent with it being only plasmid borne. However, NSS does have a full and partial copy of ISPpu12b in its chromosome (Fig. 1F). This is not surprising, considering that ISPpu12 transposes independently, in multiple copies (25), and functions in alpha-, beta-, and gammaproteobacteria (26).

Pseudomonas chloritidismutans AW-1 and Pseudomonas sp. strain PK. P. chloritidismutans AW-1’s published sequence (GenBank accession no. GQ919187) contains cld and a cytochrome $c$, which are separated from clrABDC by an insertion sequence. Similar architecture is observed in PK, which contains an insertion sequence in the exact same location (Fig. 2C). However, despite their identical positions, PK’s insertion sequence is ISPst12, while AW-1’s insertion sequence is ISPa16 (Fig. 2C). In both genomes, these insertion sequences have formed tandem duplications of TTAG. Insertion at CTAG has been previously observed for both elements (27). In PK, ISPpu12 insertion sequences again flank the genes for chlorate reduction (Fig. 2A). The ISPpu12 insertion sequences in PK are 99% identical to each other, but one (ISPpu12d) contains a notable 12-bp deletion in czd. Compared to ACDC and NSS, PK’s ISPpu12 sequences are in the opposite orientation with respect to cld and clrABDC, indicating that the composite transposons found in these bacteria were formed independently. Mapping coverage over this region (Fig. 2C) shows that the genes for chlorate reduction are present in one copy
FIG 1 Comparison of genes involved in chlorate reduction in ACDC and NSS. (A) Structure and annotation of chlorate reduction composite transposons in ACDC and NSS, including synteny with the PRI of A. suillum PS. (B) Locations of single nucleotide polymorphisms (black) in the transposase genes of ISPpu12a and ISPpu12b. (C) Composite transposon insertions in barA and nrfA of ACDC contain two flanking ISPpu12b insertion sequences. (D) Plasmids from ACDC and NSS are nearly identical and contain a chlorate reduction composite transposon flanked by isoforms ISPpu12a and ISPpu12b. T/AT and RM represent toxin/antitoxin and restriction modification systems, respectively. (E) Sequences of inverted repeats and duplications from ISPpu12a/b and ISSal1. (F) ISPpu12b inserted in several chromosomal locations in NSS.

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mbio.asm.org
A. Pseudomonas sp. PK

B. PK ISPpu12 inverted repeats

C. P. stutzeri ATCC 17588

D. PK chromosomal insertions

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(~chromosomal coverage) but that the ISPpu12d element is present in at least three other locations. We predicted these locations and confirmed them with PCR. These insertions appear to have generated duplications (Fig. 2D). In contrast, duplications at the outer edges of the composite transposon or surrounding individual insertion sequences making up the composite transposon were not found.

The left ISPpu12 insertion sequence was absent in the previously published AW-1 sequence, but the chlorate reduction genes are in the same location in the chromosome and in the middle of a conserved region in other *Pseudomonas* sequences (Fig. 2C). This allowed PCR primers to be designed outward from *dcrC* to a conserved part of *ompC* in order to finish the rest of AW-1’s sequence in this region. In contrast to PK, which contains two copies of ISPpu12, AW-1 contains only one copy and is therefore not a composite transposon (Fig. 2C).

*I. dechloratans* and *A. denitrificans* BC. *Ideonella dechloratans* was one of the first characterized chloride-reducing bacteria (28), and several fragments of genes putatively involved in chloride reduction have been sequenced, including *cl* and a conserved hypothetical gene (GenBank accession no. AJ296077.1) (29), a cytochrome *c* and molybdopterin-guanine dinucleotide biosynthesis gene (GenBank accession no. EU768872.1) (30), and *dcr-ABDC* with the insertion sequence ISIde1 (GenBank accession no. AJ566363.1) (10) (Fig. 3A). The genome sequence of *I. dechloratans* presented here expands previous sequencing efforts, identifying an *arsR* regulator, a partial methionine sulfoxide reductase gene (*msrA*), and a cupin 2 domain gene, as well as an insertion sequence that we designate ISIde2 (Fig. 3A).

Most interesting, however, is the presence of matching insertion sequences that flank this entire region in *I. dechloratans* and form a composite transposon (Fig. 3A). The insertion sequences are 99% identical to ISAAv1 (27) from *Acidovorax citrulli* AAC00-1 (GenBank accession no. AF806815), which forms a composite transposon with *s-triazine ring cleavage genes, and 99% identical to part of an insertion sequence on *Pseudomonas* sp. strain ADP’s plasmid pADP-1 (RefSeq accession no. NC_004956), which contains genes for atrazine degradation. The ISAav1 isoform in *I. dechloratans* has perfect 26-bp inverted repeats (Fig. 3B). Due to the relatively poor quality of the assembled genome (see Table S1 in the supplemental material) and highly variable coverage (Fig. 3A), it was not possible to completely resolve the location(s) of the composite transposon or deduce copy number. However, the presence of high-coverage areas (>200–versus 96-bp average coverage) suggests that it may exist in more than one copy.

*Alicyclobiplus denitrificus* BC was isolated as a benzene-degrading chlorate reducer and has been genome sequenced (31). Genes for chlorate reduction are located on a 119,718-bp plasmid, pALIDe01 (GenBank accession no. CP002450). Apart from the absence of ISIde1, the region within the *I. dechloratans* composite
transposon is remarkably similar to a region on \textit{A. denitrificans} BC pALIDE01, with just 14 single nucleotide polymorphisms in 12,546 aligned bases. A 6-bp sequence (AAAAATA) that was most likely duplicated during transposition of ISIde1 in \textit{I. dechloratans} is present only once on pALIDE01. The syntenic region (Fig. 3C, gray) includes everything carried within the \textit{I. dechloratans} composite transposon, except for the ArsR family regulator and a small hypothetical gene.

The \textit{A. denitrificans} plasmid pALIDE01 shows a history of transposition and recombination in the region surrounding chlorate reduction genes (Fig. 3C). An insertion sequence, 99.8% identical to IS1071 after reconstruction, is fragmented and found in multiple copies (Fig. 3D). The plasmid contains two copies of an ISPsy30-like transposon, which have variable 5’ regions with a conserved resolvase and transposase bound by inverted repeats (Fig. 3D). This transposon, designated ISAde1, carries a toxin/antitoxin module, as well as the \textit{ars} gene. ISAde1 flanks genes for chlorate reduction on pALIDE01 (Fig. 3C), again forming a composite architecture, although the ability of ISPsy30 transposons to mobilize large regions has not been demonstrated.

### Phyllogenetic analysis of \textit{ClrA} and \textit{Cld}

Four new genomes expand the known sequences of \textit{ClrA} and \textit{Cld} and provide insight into the evolution of chlorate reduction. Within the larger family of type II DMSO reductases, \textit{ClrA} resides in a functionally diverse clade that includes ethylbenzene dehydrogenase (EbdA), dimethylsulfide dehydrogenase (DdhA), and selenate reductase (SerA) (Fig. 4A). \textit{ClrA} proteins from \textit{I. dechloratans} and \textit{A. denitrificans} BC group with \textit{SerA} from \textit{Thauera selenatis} AX, while \textit{ClrA} proteins from PK, AW-1, ACDC, and NSS group with DdhA from \textit{Rhodovulum sulfidophilum} (Fig. 4B). In addition to the canonical \(\alpha\beta\)γ organization of genes within this clade, two additional genes appear to be conserved in the subclade containing PK, AW-1, ACDC, and NSS. The first gene is annotated as an ATPase and contains MipZ and ParA domains, and the second gene is annotated as a glycosyl transferase (Fig. 4B). The role of these genes in relation to their neighboring reductase is unknown but worthy of further investigation.

\textit{Cld} proteins in ACDC, NSS, PK, and AW-1 group phylogenetically with a betaproteobacterial clade of \textit{Cld} proteins from perchlorate reduction islands (Fig. 5A and C). In ACDC and NSS, \textit{cld} is surrounded by a \textit{napC} fragment and a small 3′ portion of \textit{pcrD} that matches the conserved gene organization of \textit{Dechloromonas aromatica} RCB and \textit{Azospira sulphii} PS (Fig. 1A) (19). In PK and AW-1, a small 3′ portion of \textit{pcrD} is present. This is highly suggestive that the direction of horizontal gene transfer was from...
FIG 4  Phylogenetic analysis of ClrA and closely related type II DMSO reductases. (A) Location of ClrA proteins within the larger type II DMSO reductase family. Asterisks signify that at least one enzyme in the clade has been shown to reduce chlorate in vitro or through genetic analysis. The black circle represents the ancestral state with proposed chlorate reductase activity. UniProt identifiers of the characterized protein are in parentheses. Conflicting biochemical data are reported in the literature for SerABC from *A. denitrificans* (44, 74). (B) Phylogenetics of ClrA and closely related enzymes. Gene neighborhoods reveal two additional conserved genes.

an RCB/PS-type PRI to a chlorate reduction composite transposon.

In *A. denitrificans* BC, *I. dechloratans*, and PK, an additional *clr*-like gene was found unassociated with chlorate reduction genes. These encode Cld-like proteins that form a unique type II subclade (Fig. 5A), with exaggerated signatures of horizontal gene transfer (Fig. 5B). In *I. dechloratans*, the *cld*-like gene is a passenger on an ISPa38-like transposon, designated ISIde3. ISPa38 transposons contain a cupin 2 domain gene and are found in the genomes of *Aeromonas caviae*, *Actinobacillus sp.* PK, *Pseudomonas metallidurans* CH34, and *Burkholderia cepacia*. The *cld*-like gene appears to have replaced a portion of the ISPa38 transposase (Fig. 5B). In *A. denitrificans* BC, the *cld*-like gene is colocated with a cupin 2 domain gene on the plasmid-borne transposon ISAde1. ISAde1 is found in *Ralstonia pickettii* 12D and 12J and in partial form in *Capriavida metallidurans* CH34 (Fig. 5B). The model tree species *Populus trichocarpa*, which can reduce perchlorate and chlorate (32), contains a *cld*-like gene that is 99.6% identical to the copy in ISAde1. The contig with the *Populus trichocarpa cld* also contains a small portion of the antitoxin gene from ISAde1, suggesting that it is bacterial in origin (Fig. 5B). PK contains a *cld*-like gene and a cupin 2 domain gene that share synteny with a *Pseudomonas stutzeri* 1A501 region comprising phage-associated, UV resistance genes (Fig. 5B).

**DISCUSSION**

Despite a series of analogous biochemical reactions, genes for chlorate and perchlorate reduction have different genomic architectures and a distinct, yet intertwined, evolutionary history. Composite transposons containing chlorate reduction genes with different flanking insertion sequences (type and ori-entation) have been identified in five chlorate-reducing bacteria. The transposon ISAde1 that surrounds *cld* and *clr* in *A. denitrificans* BC (Fig. 3C) has not been reported to mobilize genes as a composite and may not be relevant to horizontal transfer of chlorate reduction. However, ISPpu12 and ISAvl isosforms have been observed in a composite architecture previously, and the former can transpose as a composite (26).

Identification of 8-bp direct repeats at the outermost edges of the composite transposon in *nrA*, *barA*, and the plasmid (Fig. 1E) in ACDC strongly suggest that the entire element transposed into these locations. These duplications probably occurred when staggered cuts in the target site were filled in following transposon insertion (20). This, together with the fact that chlorate reduction composite transposons can be plasmid borne, provides a conceivable mechanism by which the metabolism moves horizontally. Indeed, the 99.9% nucleotide identity between plasmids in ACDC and NSS is indicative of...
Chlorate Reduction Composite Transposons

FIG 5  Phylogenetic analysis of Cld and Cld-like proteins. (A) Respiratory Cld proteins form a monophyletic clade (green) that is distinct from other Cld-like proteins found in many chlorate reducers (orange). Asterisks signify chlorite dismutase activity in vitro or through genetic analysis. Circles visually represent bootstrap support at nodes. (B) Superimposing gene neighborhoods highlights the association of cld-like genes with cupin domain genes and mobile genetic elements in a type II subclade. Cld is shown in blue at the center of the gene alignment, while the cupin 2 protein is green and usually located directly adjacent to Cld. (C) Cld proteins from chlorate-reducing strains ACDC, NSS, PK, and AW-1 group with beta-proteobacterial Cld proteins from PS and RCB.
In PK, the promoter is on the opposite strand and could potentially activate genes by providing full or partial promoters (34). In ACDC, studies that assume a single stable locus.

...tions, the presence of the composite transposon becomes a metabolic burden and provides a selective advantage to cells that have lost or silenced the metabolism. Composite transposons can also increase in copy number (Fig. 1C) and confound transcriptional studies that assume a single stable locus.

Insertion sequences can activate transcription of neighboring genes by providing full or partial promoters (34). In ACDC, a −35/−10 promoter in ISPpu12 is located upstream of clrABDC. In PK, the promoter is on the opposite strand and could potentially drive clr transcription. It has been suggested that this promoter is active and constitutive in P. putida (25). Insertion sequences are also located between clr and cld in PK, AW-1, and I. dechloratans and could serve to change transcription of surrounding genes. While we cannot rule out the possibility that the repeated localization of insertion sequences between clr and cld is random, it is suggestive either of a physiological role for these insertion sequences or of the historical mechanism of colocalization of clr with clrABDC.

With the exception of A. denitrificans BC, most chlorate reducers are unable to reduce nitrate (AW-1 [35], ASK-1 [36], PDA [37], I. dechloratans [28], and NSS [unpublished data]). I. dechloratans was reported to lose the ability to reduce nitrate after cultivation on chlorate (28), and AW-1 has been reported to regain the ability to denitrify after repeated aerobic subculturing in the presence of nitrate (38). We have identified insertions in the nitrate reductase (nrfA) in ACDC and a putative nitrate-responsive histidine kinase (narX) in PK but have yet to find a genetic basis for NSS’s inability to reduce nitrate, given that it contains genes for a complete denitrification pathway (nar, nir, nor, and nos). The ability of the nitrate reductase NarGHI to reduce chlorate to toxic chloride has been widely reported (39–41). Early experiments exploited this to isolate chlorate-resistant cells with mutations in parts of the nitrate reduction pathway. This identified the nitrate reductase, as well as genes for Mo cofactor biosynthesis, molybdate transport, and nitrate regulation (42). Given this, it is not surprising that parts of nitrate reduction pathways are inactivated in some chlorate reducers, but more work is needed to fully appreciate the interplay between these metabolisms.

Perchlorate, despite having a high redox potential, has a large activation energy that slows inadvertent reduction by metals and enzymes (43). In contrast, chlorate will react abiotically with Fe(II) or Mn(II), and the evolution of an enzyme that overcomes the activation barrier for chlorate was conceivably less difficult than for perchlorate. Many related enzymes, including PcrA (9), NarG (39, 40), and SerA (44), can reduce chlorate in vitro (Fig. 4A; starred clades contain at least one characterized enzyme with chlorate reductase activity). In the case of DdhA, evolution to oxidize a hydrophobic substrate may have precluded turnover of chlorate (45). The most parsimonious hypothesis consistent with these phylogenetic data is that enzymes in this clade evolved from an ancestral enzyme capable of chlorate reduction. If this is correct, the key step in the evolution of chlorate reduction was the presence of a chlorite detoxification system.

Type I Cld proteins involved in respiratory perchlorate and chlorate reduction are monophyletic (Fig. 5A) (46) but, based on incongruence between protein and species trees, are predicted to have undergone horizontal gene transfer (46, 47). We provide evidence that cld found in four chlorate reduction composite transposons originated from a PRI, based on the transfer of small PRI-specific regions surrounding cld, including part of pcrD (ACDC, NSS, PK, and AW-1) (Fig. 5C) and in some cases a napC homolog (ACDC and NSS) (Fig. 1A). Given that a PRI cld has been coopted for chlorate reduction, why has an active type II chlorite dismutase (46) (Fig. 5A) yet to be associated with a respiratory metabolism? One barrier is the lack of a signal sequence and therefore cytoplasmic localization, which would not protect cells from chlorite produced periplasmically. Another difference is structural: the type II Cld from Nitrobacter winogradskyi is dimeric (48) compared to the tetrameric (49, 50) or pentameric/hexameric (51, 52) type I Cld proteins, but the implications of this difference are unknown. It is also possible that the diversity of respiratory Cld has not been fully surveyed.

In addition to type I Cld, several chlorate reducers contain Cld-like proteins located elsewhere on the chromosome that form a subclade within the type II lineage (Fig. 5A, orange). This subclade contains a set of cld-like genes that are part of transposons (ISPa38-like, ISPsy30-like, and IS26 composite), or associated with phage-related genes, and are therefore presumably highly mobile (Fig. 5B). In all but Acinetobacter baumannii OIFC074, a cupin 2 domain gene is colocated with cld; this occurs in multiple configurations and on different transposons (Fig. 5B). Colocalization of type I cld and cupin genes also occurs in some chlorate reduction composite transposons (Fig. 3A) and PRIs, and as a result, we hypothesize a functional connection between these genes.

Comparative analysis of composite transposons identified the core metabolic genes and several additional genes that may be important for chlorate reduction. These are relatively scarce compared to those in perchlorate reduction islands, suggesting that this mobile metabolism often relies on endogenous systems for electron transport, cofactor biosynthesis, and regulation. At the least, a connection to the host’s electron transport chain must be established. By analogy to the closely related enzymes SerABC (53) and DdhABC (54), and as previously proposed (55), it is possible that CldABC connects to a quinol oxidase via a cytochrome c. ACDC, NSS, PK, and AW-1 all contain a cytochrome c next to cld that is a possible candidate. However, in I. dechloratans, the cytochrome c in the chlorate reduction composite transposon does not donate electrons to CldABC in vitro, suggesting that it does not provide the necessary link to the electron transport chain (30). Another cytochrome c in I. dechloratans has been shown to donate electrons to CrABC in vitro (56), and we identified its sequence (see Fig. S2 in the supplemental material) based on the fragment reported in the literature (55). A quick search (57) for similar proteins gave NirM, the electron donor for nitrite reductase NirS (58), as a top hit.

Regulation of chlorate reduction with respect to other electron
acceptors is not well understood. Lack of regulatory genes in the composite transposons of ACDC, NSS, PK, AW-1, and BC implies that regulation may be absent or controlled by a chromosomal system. In *L. dechlororans*, an ArsR family regulator was identified that may be involved in the observed increase of *cld* or *clrA* transcription under anaerobic conditions in the presence of chlorate (59). Although this family of regulators is usually associated with sensing metals, evidence exists for nonmetal signals (60).

The discovery of multiple, independently formed, chlorate reduction composite transposons advances our understanding of this metabolism at the gene level and supports the idea that chlorate reduction is a highly mobile metabolism. Flanking insertion sequences help define a set of genes that move with *cld*ABDC and *cld*, and although some of these genes may be passengers that were captured as the composite transposons formed, others are likely to have roles in chlorate reduction. We have identified these genes by comparative analysis and hypothesized functions for several. The next step is to develop a genetic system in a chlorate reducer and make clean deletions to test these predictions.

**MATERIALS AND METHODS**

**Genome sequencing.** Whole-genome shotgun sequencing and initial assembly of four chlorate-reducing bacteria, *Idonella dechlororans*, *Pseudomonas* sp. strain PK, *Dechloromonas chlorophurus* NSS, and *Shewanella algae* ACDC, were completed by Eureka Genomics (Hercules, CA). A summary of this sequencing effort is provided in Table S1, and a summary of genes internal to chlorate reducing composite transposons is provided in Table S3. Annotation of the genomes was publically available through IMG (http://img.jgi.doe.gov/cgi-bin/w/main.cgi). A summary of genes internal to chlorate reduction composite transposons is provided in Table S3. Annotation of the manually assembled plasmid from ACDC and NSS was performed using RAST (62).

**Improving draft assemblies.** Bowtie (63) was used to map reads (Bowtie -q -n 3 -i 60 -e 200 – best) to assembled genomes. Multiple hits were not allowed in the Bowtie mapping stage. IMAGE (24) was used to extend assembled contigs when possible, and extensions were blasted against all other contigs to search for potential linkages (BLASTn -word_size 10) (64). It should be noted that not all extensions were valid; some were later determined to be erroneous after finishing PCRs were completed. Linkages were used to create a contig graph surrounding chlorate reduction genes. This graph was further simplified using coverage to understand the relative abundance of duplicated regions, which was important for the identification of composite transposons. This workflow is further detailed in Text S1 and Fig. S3 in the supplemental material. PCR and sequencing were performed to confirm linkages. A list of finishing primers is found in Table S4. To further support predicted *in silico* linkages, reads were remapped to contig junctions and coverage was determined with samtools mpileup (65).

**Gene visualization.** Visualization of genes was performed using python scripts and the GenomeDiagram package (66). Additional modifications to diagrams were performed in Adobe Illustrator. ApE was used for viewing and comparing nucleotide sequences and for manually annotating features. Mapped reads and sequenced PCR products were visualized by overlaying them on assembled contigs. Horizontal bars represent PCR products, with black indicating the portion that was Sanger sequenced. Inverted repeats are shown as black vertical lines at the outer edges of insertion sequences. Syntenic regions of high nucleotide identity are highlighted in gray.

**Phylogenetics.** Phylogenetics was performed using a compute cluster running CentOS 5.6 at UC Berkeley’s QB3 Computational Genomics Resource Laboratory. Proteins are labeled in figures with an identification code reflecting their source: uniprot = UniProtID, refseq = NCBI ReSeq ID, gi = NCBI GI number, gb = GenBank locus or accession number, and img = IMG GeneID. The alpha subunit of chlorate reductase (ClrA) from ACDC was used as the query to search IMG and NCBI (nr database) for closely related proteins (BLAST-P). After duplicates were removed, the top 100 sequences were added to prealigned DMSO reductase type II family proteins (Pfam 00384 and 17524 sequences; http://pfam.sanger.ac.uk/, accessed 4 December 2012). CD-HIT (67) was run with a clustering threshold of 0.7, the reduced protein alignment was trimmed with Gblocks (b1 = N/2+1-b2 = N/2+1-b3 = N/2-b4 = 2 -b5 = h, where N is the number of sequences) (68), and the protein phylogeny was reconstructed with FastTree (69) in order to obtain an outgroup basal to the clades of interest. Sequences exterior to this outgroup were discarded, and the remaining sequences were realigned and trimmed. The best amino acid substitution model was determined with ProtTest (70) to be MT-MAM. Subsequent phylogenetic analysis was performed using RAxML with 500 bootstraps (71). The same procedure was followed when building a phylogeny for the subclade containing ClrA. Chlorite dismutase (Cld) sequences were obtained with a PSI-BLAST against NCBI (three iterations, e = 10^-3) using ACDC’s type I Cld as the starting query. These sequences were combined with the top 100 hits from a BLAST-P search of both IMG and NCBI (nr) that used the *A. denitrificans* Cld-like protein as a query. The same workflow as above was used to select an appropriate outgroup and reconstruct the phylogeny with RAxML. 16S RNA gene sequences from perchlorate- and chlorate-reducing bacterial isolates were obtained from NCBI and newly sequenced genomes. Three nearest-neighbor matches were found and aligned using Ribosomal Database Project tools (72). Phylogenetic analysis was performed with MrBayes 3.2 (73) until the average standard deviation of the split frequencies was less than 0.01. Posterior probabilities were estimated after discarding the first 25% of samples from the cold chain.

**SUPPLEMENTAL MATERIAL**

Supplemental material for this article may be found at http://mbio.asm.org/lookup/suppl/doi:10.1128/mBio.00379-13/-/DCSupplemental.

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