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Involvement of β -carbonic anhydrase (β -CA) genes in bacterial genomic islands and horizontal transfer to protists

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ABSTRACT

Genomic islands (GIs) are a type of mobile genetic element (MGE) that are present in bacterial chromosomes. They consist of a cluster of genes which produce proteins that contribute to a variety of functions, including, but not limited to, regulation of cell metabolism, anti-microbial resistance, pathogenicity, virulence, and resistance to heavy metals. The genes carried in MGEs can be used as a trait reservoir in times of adversity. Transfer of genes using MGEs, occurring outside of reproduction, is called horizontal gene transfer (HGT). Previous literature has shown that numerous HGT events have occurred through endosymbiosis between prokaryotes and eukaryotes.

Beta carbonic anhydrase (β -CA) enzymes play a critical role in the biochemical pathways of many prokaryotes and eukaryotes. We have previously suggested horizontal transfer of β -CA genes from plasmids of some prokaryotic endosymbionts to their protozoan hosts. In this study, we set out to identify β -CA genes that might have transferred between prokaryotic and protist species through HGT in GIs. Therefore, we investigated prokaryotic chromosomes containing β -CA-encoding GIs and utilized multiple bioinformatics tools to reveal the distinct movements of β -CA genes among a wide variety of organisms. Our results identify the presence of β -CA genes in GIs of several medically and industrially relevant bacterial species, and phylogenetic analyses reveal multiple cases of likely horizontal transfer of β -CA genes from GIs of ancestral prokaryotes to protists.

IMPORTANCE

The evolutionary process is mediated by mobile genetic elements (MGEs), such as genomic islands (GIs). A gene or set of genes in the GIs are exchanged between and within various species through horizontal gene transfer (HGT). Based on the crucial role that GIs can play in bacterial survival and proliferation, they were introduced as the environmental- and pathogen-associated factors. Carbonic anhydrases (CAs) are involved in many critical biochemical pathways, such as regulation of pH homeostasis and electrolyte transfer. Among the six evolutionary families of CAs, β -CA gene sequences are present in many bacterial species, which can be horizontally transferred to protists during evolution. This study shows for the first time the

involvement of bacterial β -CA gene sequences in the GIs, and suggests their horizontal transfer to protists during evolution.

KEYWORDS

β -carbonic anhydrase; Evolution; Genomic island; Mobile genetic element

Horizontal Gene Transfer (HGT) is an evolutionary phenomenon by which a gene, or set of genes, are exchanged between and within various species. This makes HGT unique compared with other evolutionary processes, such as gene duplication, mutation, and sexual reproduction. While a heritable HGT in eukaryotes entails entrance of a foreign gene to the nucleus of the germ cell and successful insertion to chromatin packed DNA, HGT has multiple pathways in prokaryotic species. This evolutionary process is mediated by mobile DNA or mobile genetic elements (MGEs), which can include: genomic islands (GIs), plasmids, transposons, retrotransposons, and prophages (1-6). During HGT, selfish “parasitic” elements are often associated with toxin resistance genes, metabolic genes, virulence factors, and a wide range of secreted factors. The acquisition of a useful gene repertoire could offset the cost of maintaining and transferring a large selfish element, such as a conjugal plasmid (7). Transformation, conjugation, and transduction are each distinct methods of HGT in prokaryotes.

Varieties of important genes are transferred between prokaryotes, or from prokaryotes to eukaryotes, through HGT (8), including those for virulence factors, antibiotic resistance, and toxins (1-3). In 1990, some clusters of virulence genes, which transfer through HGT, were identified in *Escherichia coli* and described as pathogenicity islands (PAIs) (9). Later GIs were defined as any cluster of genes (10–200 kb) that has been acquired by HGT (10). GIs represent a part of a cell’s chromosome, recognized as discrete DNA segments, and can differ between closely related strains. Different GI families have been recognized on the basis of sequence and functional homologies by GI prediction tools (11). The nucleotide sequence length of GIs is >10 Kb, while it is <10 Kb for smaller genomic islets (12).

Interest in GIs has increased commensurately with developing knowledge of their role in bacterial survival and proliferation. The common environmental- and pathogen-associated virulence factors found disproportionately in GIs tend to serve functions. For example, the pathogenicity role of β -CA has been approved in *Pseudomonas aeruginosa* (13) and the critical role of β -CA in detoxification of cyanate by providing bicarbonate for cyanase enzyme has been shown in *Pseudomonas pseudoalcaligenes* (14).

However, clustered regularly interspaced short palindromic repeats (CRISPRs), used by bacteria in defense against insertion of phage DNA, are also found overrepresented in GIs (15).

The presence of GIs have been studied using various computational biology methods. There are two main methods for prediction of GIs, including: (1) evaluation of sequence compositions, using tools such as SIGI-HMM (16), IslandPath-DIMOB (17), PAI-IDA (18), and Centroid (19), and (2) application of comparative genomics, such as BLAST homology search and whole-genome sequence alignment. Among the sequence analysis methods, SIGI-HMM and IslandPath-DIMOB have shown the highest overall accuracy (15). Two computational methods for prediction of GIs based on comparative genomics, include IslandPick (20) and MobilomeFINDER (21). The latter method focuses on identification of the islands associated with *tRNA* genes. However, not all GIs use *tRNA* genes as insertion sites, which thus limits the usage of MobilomeFINDER compared to the IslandPick method (22). Prediction based on IslandPick is provided at the IslandViewer 4 database [<http://www.pathogenomics.sfu.ca/islandviewer/>] (23). The IslandViewer server combines the three most accurate GI prediction methods into a single analysis: IslandPath-DIMOB, SIGI-HMM, and IslandPick.

Carbonic anhydrases (CAs) are ubiquitous metalloenzymes, which are categorized into seven gene families, including α , β , γ , δ , ζ , η , and θ (24-27). CAs are involved in many important biochemical pathways including pH homeostasis, electrolyte transfer, transport of CO₂ and bicarbonate between metabolizing tissues, and some biosynthetic processes (28-31). Many ancient putative β -CAs have been discovered in protozoans, rotifers, sea louses, molluscs, starlet sea anemones, purple sea urchins, arthropods, nematodes, and trematodes (32-34), as well as in prokaryotes and some eukaryotes, such as fungi, algae, and plants (35). Notably, β -CA gene sequences are present in the genomes of most living organisms except vertebrates (33, 34). β -CAs are considered to be crucial metabolic enzymes (32, 36, 37). They act as virulence factors for various bacterial, fungal and parasitic species, such as *Pseudomonas aeruginosa* (13), *Cryptococcus neoformans* (38), and *Toxoplasma gondii* (39), so β -CAs develop a cascade leading to the production of infectious spores in *C. neoformans*, prepare the adaptation of *P. aeruginosa* to low CO₂ condition through

different organization of three β -CA genes, and play the role in rhoptry biogenesis and formation of parasitophorous vacuole in *T. gondii*. β -CA is a vital enzyme for fertility of female insects (*Drosophila melanogaster*) (36) and therefore, β -CAs are attractive targets for inhibition studies in insects and pests. There is active ongoing research in this field focusing on inhibition of β -CAs in important organisms; for example, application of sulfonamide and sulfamate for inhibition of β -CA from *Helicobacter pylori* (40), aromatic carboxylates for inhibition of β -CA from *Candida albicans* (41), sulfonamides for inhibition of β -CA from *Ascaris lumbricoides* (42), and sulfonamides for inhibition of β -CA from malaria mosquito *Anopheles gambiae* (43). CA inhibition studies have been mainly performed *in vitro*, and only a few *in vivo* studies have been carried out on parasitic infectious diseases (44-46).

Here we have studied the importance of HGT and β -CA gene exchange between bacterial GIs and protists genomes. We propose that GIs play a crucial role in horizontal transfer of β -CA genes from prokaryotes to protists.

RESULTS

β -CA genes are located in many genomic islands

Our comparative analysis of the GI annotations presented in the IslandViewer 4 database and NCBI genome annotations, allowed us to identify a total of 272 instances of β -CA genes in bacterial GIs (Table S1). In study of all strains, nucleotides in β -CA genes are 3.81x more likely to occur in GIs than is expected by chance.

Identification of β -CAs from prokaryotes and protists

A multiple sequence alignment (MSA) was created for 86 amino acid residues of 25 prokaryote and protist β -CA protein sequences. The alignment revealed that all β -CA protein sequences contain the first (CXDXR; C: Cysteine, D: Aspartic acid, R: Arginine, and X: any amino acid) and second (HXXC; H: Histidine, C: Cysteine, and X: any amino acid) highly conserved motifs, which are characteristic of a β -CA protein (Fig. 1).

Using the data from the IslandViewer 4 IslandPath-DIMOB webserver and corresponding NCBI genome annotations, 272 β -CA genes were identified inside of prokaryote GIs (Table S1) (e.g. β -CA-encoding GI from *Methylibium petroleiphilum* (strain PM1)).

Table S1. Prokaryotic GIs containing β -CA genes.

Phylogenetic analysis

The result of the phylogenetic analysis is presented as a circular tree and divisions of interest are shown in three different clades: A, B, and C (Fig. 2). Partitioning delineates regions where β -CA genes appear to have a common ancestor in prokaryotes and protists.

In clade A, the β -CA gene of *Trichomonas vaginalis* and *Paulinella chromatophora* has a common ancestor with β -CA genes from prokaryotic GIs. In Clade B, there is a β -CA gene from a prokaryotic GI and two β -CA genes from the protist *Acanthamoeba castellanii*. Clade C includes β -CA genes from 27 bacterial GIs and 14 protists including *A. castellanii*, *Capsaspora owczarzaki*, *Dictyostelium discoideum*, *D. fasciculatum*, *D. purpureum*, *Leishmania donovani*, *L. panamensis*, *Leptomonas pyrrocoris*, *Phaeodactylum tricornutum*, *Phytophthora infestans*, *Polysphondylium pallidum*, *Saprolegnia diclina*, *Tetrahymena thermophila*, and *Trypanosoma grayi*. We did not identify any definite bacterial common ancestor with β -CA genes from protists *Entamoeba invadens*, *E. nuttalli*, and *Galdieria sulphuraria*.

Sequence conservation analysis for HGT

In order to evaluate the hypothesis of HGT between prokaryotes and eukaryotes within Clade C, the sequence conservation among the Clade C proteins were compared to the rest of the phylogenetic tree. First, Clade C protein sequences were aligned using Clustal Omega and the residues fully conserved within Clade C were identified (14 residues). Then, all proteins within the phylogenetic tree outside Clade C, except protist sequences, were aligned using Clustal Omega (Fig. S1). The resulting MSA was analyzed using program Consurf for sequence conservation (Fig. 3). We then inspected the conservation of those 14 fully-conserved residues within Clade C for their conservation in the large group of CA sequences. Among those, 7 residues were highly conserved (conservation score 9) and 4 were well conserved (conservation score 7-

8). However, 3 of the residues showed an average of low conservation (conservation score 3-5). Therefore, conservation of those 3 residues could be considered as a possible result of HGT. These three conserved residues (Leu21, Gly71 and Gly117) of the homology modeled of β -CA from *T. vaginalis* (A2ENQ8) were shown in Fig. 4.

FIG S1 Multiple sequence alignment (MSA) for all proteins within the phylogenetic tree outside Clade C (Fig. 2), except protist sequences, were aligned using Clustal Omega.

Exon count for β -CA genes from protists

Single exon structure of a protist gene would provide some additional support for the hypothesis that this particular gene could be of prokaryotic origin. The exon count analysis revealed that the β -CA genes of certain protists have indeed a single exon, while many other β -CA genes have multiple exons. The single-exonic β -CA genes of protists have shown in Table 1.

DISCUSSION

Our previous phylogenetic analysis has suggested that β -CA genes have crossed species boundaries on multiple occasions (8). The present identification of β -CA genes within bacterial GIs also strongly suggests that prokaryotic β -CA genes have been horizontally transferred between and within different species (47). Specifically, we see what appears to be a very clear case of HGT of a β -CA from prokaryotic GIs to protists *T. vaginalis* and *P. chromatophora* (clade A), *A. castellanii* (clade B), and *Dictyostelium* sp., *P. pallidum*, *T. thermophila* (NCBI IDs: XP_001009612.1, XP_001013978.2, XP_001022390.2), *Leishmania* sp., *L. pyrrocoris*, and *T. grayi* (clade C). Also, our phylogenetic analysis reveals that β -CA genes from GIs of *Bacillus thuringiensis* and *Psychrosinus fermentans* have common ancestors with *T. vaginalis* (clade A) and *A. castellanii* (clade B), respectively. In addition, a β -CA gene from a GI of *Rahnella aquatilis* shows a common ancestor with *Dictyostelium* sp. and *P. pallidum*. In several cases we observe clustering of multiple β -CA genes from the same protist species. When these cluster immediately together we believe the most likely explanation is gene duplication after HGT. In the case where we observe two distinct clusters of

paralogs from the same protist, such as with *T. thermophila* in Clade A, it is possible that there have been multiple duplication events after HGT, or separate cases of HGT.

A previous study has shown that β -CA genes in protists exist as single or multiple exon chromosomal genes, while in metazoans these genes exist only as multiple exon chromosomal genes (48). Single exon β -CA genes can be found in most of our candidate HGT species, *Entamoeba* sp., *Leishmania* sp., *L. pyrrocoris*, *P. chromatophora*, *Phytophthora infestans* (XP_002909250.1), *T. thermophila* (XP_001009111.1), *T. vaginalis* (NCBI ID: XP_001317907.1), and *T. grayi*; while multiple exon β -CA genes can be found in the other protists. The single exon structure of some β -CA genes of protists thus suggests that they are closely associated to prokaryotic β -CAs. Therefore, it seems that β -CA genes with prokaryotic GIs origins have integrated into stable chromosomal loci in genomes of protists without association between GIs and β -CA genes of protists. Due to the large and complexity of eukaryotic genomes and heterogeneous chromosomes leading to high rates of false-positive results, the horizontal transfer of GIs to the eukaryotic genomes is in a halo of ambiguity and largely unexplored (49). Currently, identification studies of horizontally transferred genes to eukaryotes are performed through comparative analyses than experimental methods to show GIs in the genome of the eukaryotes. Therefore, further studies are needed to design the databases for identification of eukaryotic GIs.

Our studies have revealed that β -CA protein has, in some cases, potentially evolved into a virulence factor for some pathogenic bacteria, such as *B. pseudomallei* (50). We have identified a significant number of β -CA genes which reside in GIs in bacterial species, many of which are known to be pathogenic. These β -CA genes occur inside of GIs at a significantly higher rate than expected by chance, implying some function. Known virulence factors can be influenced by attenuators or RNA-based regulatory strategies, which lead to premature termination of transcription (51). Based on the lack of β -CAs in vertebrates, these proteins can be considered potential targets for anti-parasitic drugs. On the other hand, due to the presence of β -CA in *M. petroleiphilum*, it is suggested that this enzyme plays a major role in a CO₂-concentrating-mechanism (CCM) in carboxysomes through use of methyl tert-butyl ether (MTBE) as the sole source of carbon. The β -

CAs from other extremophilic bacteria (Table S1), such as *Halothermothrix orenii* (52), *Acidithiobacillus caldus* (53), *Thioalkalivibrio nitratireducens* (54), and *Thiomicrospira crunogena* (55) may play critical metabolic roles through carboxysome or non-carboxysome-associated mechanisms.

The GIs in some bacterial species, such as *Pseudomonas aeruginosa*, can lead to emergence of strains resistant to various antibiotics (56). It was demonstrated that *P. aeruginosa* (isolate ST235) contains Tn6162 and Tn6163 in GI1 and GI2, respectively, which function together as multiple antibiotic-resistant cassettes. An environmental study showed that *Thiomonas* sp. is able to withstand the extreme conditions of acid mine drainage (57). The comparison between the genomes of *T. arsenitoxydans* (strain 3As), *T. intermedia* (strain K12), and *Thiomonas* sp. (strain CB2) identified over 20 GIs occurring through various rearrangements containing arsenite resistance and oxidation genes, leading to divergent resistance to arsenic-rich environments.

Conclusions. A GI is a continuous genomic region which arises through HGT and can contain tens to hundreds of genes. We have previously identified cases of horizontal transfer of β -CA genes from plasmids of some prokaryotic endosymbionts to their protozoan hosts (48). The present results support the idea that β -CA genes in protists and modern eukaryotes originated by HGT from ancestral prokaryotic GIs, along with other facilitators, such as transposase and integrase. Using phylogenetics and homology modeling, we suggest that the close sequence similarity of CA genes in hosts and endosymbionts was due to HGT and not convergent evolution (48).

Further studies will be needed to identify the origin of β -CA genes in ancestral and metazoan species. Even though our results suggest that β -CA genes are overrepresented in GIs compared to the rest of the genome, no studies have yet been reported on whether β -CA genes are overrepresented there compared to other metabolic genes.

MATERIALS AND METHODS

Identification of CA proteins in genomic islands

A total of 110,913 genomic island annotations for 6,348 complete bacterial and archaeal strains were retrieved from the IslandViewer 4 database. RefSeq assembly annotations were available for 6,238 strains, which were downloaded in bulk from the NCBI assembly server (<https://www.ncbi.nlm.nih.gov/assembly>); annotations only available GenBank and updated annotation versions not corresponding to the current IslandViewer release were not retrieved. Using custom Python scripts, assembly IDs were retrieved for all IslandViewer genome accession IDs, and gene annotations compared with the GI locations. All CA genes which occur within the IslandViewer defined GI locations were kept for further analysis. To determine average overrepresentation, for each examined genome the number of CA gene nucleotides (nt) overlapping any GI was compared to the number expected by chance alone. Expected overlap was defined as the sum of lengths of all CAs in the genome multiplied by the sum of lengths of all GIs in the genome divided by the length of the genome.

Identification of β -CAs from prokaryotes and protists

After detection of GIs-containing β -CA genes in the IslandViewer version 4 database, we then collected prokaryotic β -CAs locating in GIs, and β -CA from *Klebsiella pneumoniae* subsp. *pneumoniae* (NCBI protein ID: WP_019705531.1), five β -CA protein sequences equally from both gram-negative and gram-positive bacteria (Table 2), and protist β -CAs (Table 3) to perform a multiple sequence alignment (MSA) analysis. We used the β -CA protein sequence from *Klebsiella pneumoniae* subsp. *pneumoniae* (NCBI protein ID: WP_019705531.1) as a query from prokaryotic species for the MSA analysis. All β -CA protein sequences from protists used in the analysis are described in Table 3. Also, we used the β -CA protein sequence from *A. castellanii* (XP_004344666.1) as the query from protists in the Basic Local Alignment Search Tool for proteins (blastp) from NCBI database (https://blast.ncbi.nlm.nih.gov/Blast.cgi?PROGRAM=blastp&PAGE_TYPE=BlastSearch&LINK_LOC=blasthome) through running different phyla of protists including Stramenopiles, Alveolata, Rhizaria, Excavata, Amoebozoa, Hacrobia, Apusozoa, and Opisthokonta in the choosing search set panel. Some protists contain more than one β -CA protein sequence, in which case we used only one as a representative sequence in the

MSA analysis. In total 25 β -CA protein sequences of prokaryotes and protists (86 amino acid residues, starting three amino acid residues prior to first highly conserved motif; CXDXR) were used to compute an MSA using the Clustal Omega. The results were visualized in JalView [<http://www.jalview.org/>] (58).

Identification of β -CA gene sequences located in prokaryotic GIs was performed using the IslandViewer version 4 database. This webtool provides the ability to draw main circular chromosomes of defined prokaryotes containing GIs, as well as search for β -CA gene sequences.

Phylogenetic analysis

The β -CAs identified to reside in bacterial GIs, using annotations from the IslandViewer database, were clustered to 90% similarity centroids with the "cluster_fast" algorithm of the search tool (59) in order to reduce the number of sequences for phylogenetic analysis. Similarly, a set of 35 protist β -CAs were clustered to 90% similarity. The resulting reduced set of 122 prokaryote β -CAs found within GIs and 35 protist β -CAs were aligned using Clustal Omega. Model testing was performed to identify the best evolutionary model for analysis of the target sequences using ModelFinder (60). A maximum likelihood phylogenetic analysis was performed using the IQTree software (61, 62), with parameters set to "-alrt 100000 -bb 100000 -nt AUTO -m LG+R7" and all other options run as default. A consensus tree was generated from the 100,000 bootstrap replicates, with a final log-likelihood value of -37626.11. The tree was then visualized using the ETE Toolkit Python library (63).

Sequence conservation analysis for HGT

In order to analyze sequence conservation among the β -CA proteins, their sequences were aligned using the Clustal Omega and the resulting MSA was then analyzed using the ConSurf Server [<http://consurf.tau.ac.il/>] (64).

Exon count for β -CA genes from protists

In order to count the exons of β -CA genes from protists, we used the NCBI gene server (<https://www.ncbi.nlm.nih.gov/gene/>) (65). In this feature, a summary of a specific gene including gene

type, symbol and description, locus tag, RNA name, RefSeq status, organism lineage, and genomic context (exon count) are presented.

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Authors' contributions: All authors participated in the design of the study. RZE carried out the search and collection of relevant prokaryotic and eukaryotic species, identification of β -CAs, and conservation analysis. RZE created the MSAs. HRB identified the β -CA genes in GIs, made protein sequence corrections and predictions, and performed the phylogenetic analysis. VPH performed sequence conservation analysis and protein modeling. RZE drafted the first version of the manuscript. All authors participated in writing, read and approved the final manuscript.

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TABLE 1 The single exon β -CA genes of protists.

No.	Protist species	NCBI IDs	Gene name
1	<i>Entamoeba</i> sp.	XP_004183626.1	EIN_065450
		XP_008860421.1	ENU1_204230
2	<i>Leishmania</i> sp.	XP_003858369.1	LDBPK_060630
		XP_010703940.1	LPMP_060590
3	<i>Leptomonas pyrrocoris</i>	XP_015662104.1	ABB37_01925
		XP_015662099.1	ABB37_01923
4	<i>Paulinella chromatophora</i>	YP_002049530.1	PCC_0911
5	<i>Phytophthora infestans</i>	XP_002909256.1	PITG_00682
		XP_002909250.1	PITG_00674
6	<i>Tetrahymena thermophila</i>	XP_001009111.1	TTHERM_00263620
7	<i>Trichomonas vaginalis</i>	XP_001317907.1	TVAG_005270
8	<i>Trypanosoma grayi</i>	XP_009310034.1	DQ04_02331000

TABLE 2 β -CA protein sequences from bacterial species.

No.	Gram-staining	Bacterial species	NCBI IDs
1	Gram-negative	<i>Klebsiella pneumoniae</i>	WP_019705531.1
2		<i>Brucella abortus</i>	WP_002965854.1
3		<i>Yersinia enterocolitica</i>	WP_005165125.1
4		<i>Bordetella parapertussis</i>	YP_006895229.1
5		<i>Pseudomonas stutzeri</i>	WP_011914306.1
6	Gram-positive	<i>Streptomyces</i> sp.	WP_015579823.1
7		<i>Bifidobacterium angulatum</i>	WP_003825226.1
8		<i>Pseudonocardia</i> sp.	WP_060712833.1
9		<i>Desulfocapsa sulfexigens</i>	WP_015403686.1
10		<i>Arthrobacter alpinus</i>	WP_062006860.1

TABLE 3 β -CA protein sequences from protists.

No.	Protist species	NCBI IDs
1	<i>Acanthamoeba castellanii</i>	XP_004344666.1, XP_004335990.1, XP_004337607.1
2	<i>Capsaspora owczarzaki</i>	XP_004342925.1, XP_4349240.1
3	<i>Dictyostelium</i> sp.	XP_646739.1, XP_644170.1, XP_003283430.1, XP_004361116.1
4	<i>Entamoeba</i> sp.	XP_004183626.1, XP_008860421.1
5	<i>Galdieria sulphuraria</i>	XP_005703553.1
6	<i>Leishmania</i> sp.	XP_003858369.1, XP_010703940.1
7	<i>Leptomonas pyrrocoris</i>	XP_015662104.1, XP_015662099.1
8	<i>Paulinella chromatophora</i>	YP_002049530.1
9	<i>Phaeodactylum tricornutum</i>	XP_002176594.1
10	<i>Phytophthora infestans</i>	XP_002909256.1, XP_002909250.1, XP_002909249.1
11	<i>Polysphondylium pallidum</i>	XP_020436034.1
12	<i>Saprolegnia diclina</i>	XP_008607403.1, XP_008604330.1
13	<i>Tetrahymena thermophila</i>	XP_001009617.1, XP_001009612.1, XP_001009111.1, XP_001022390.2, XP_001009116.2, XP_001009616.1, XP_976601.1, XP_001013978.2
14	<i>Trichomonas vaginalis</i>	XP_001317907.1, XP_001579768.1
15	<i>Trypanosoma grayi</i>	XP_009310034.1

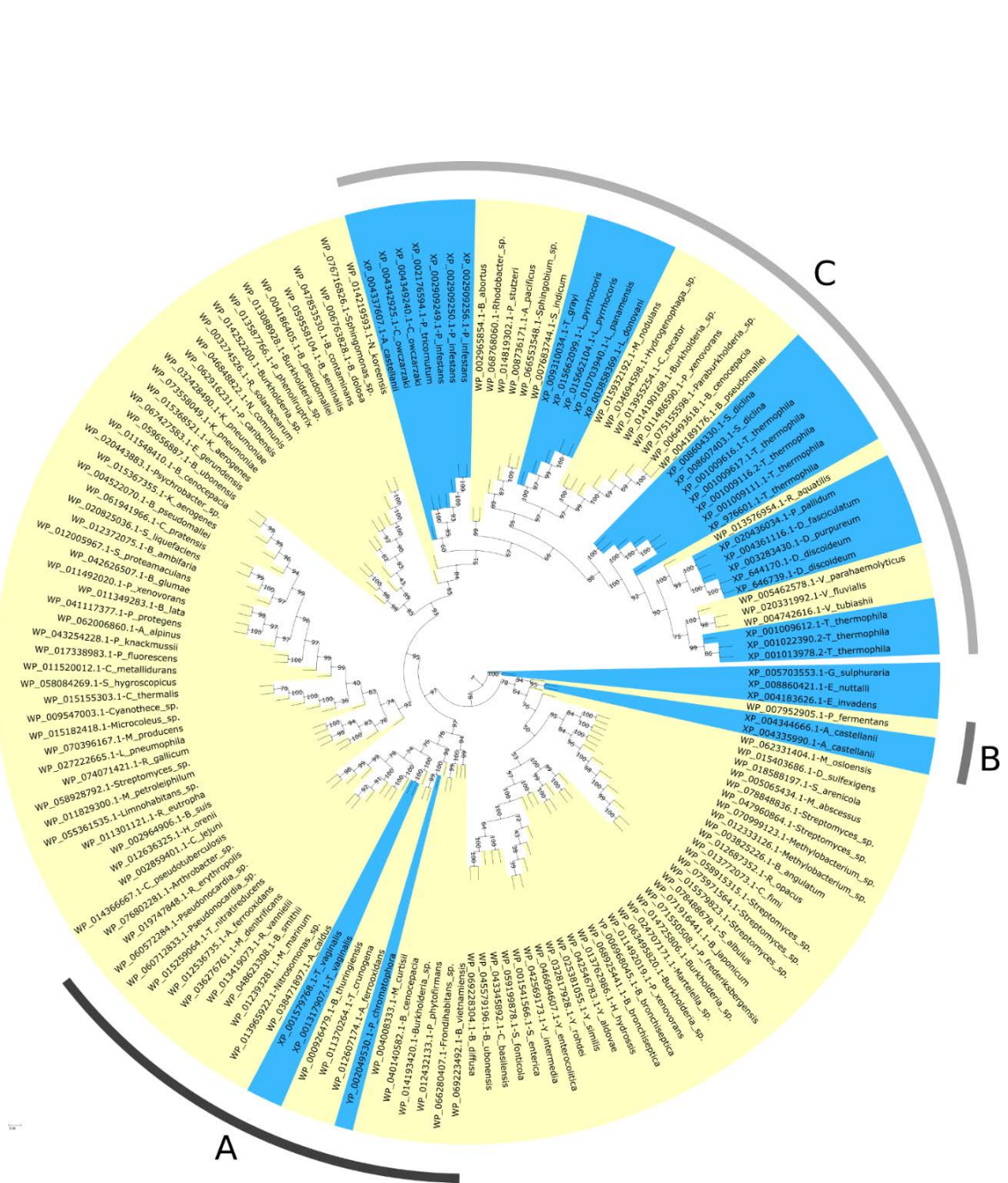


FIG 2 Phylogenetic analysis of β -CAs from prokaryotes and protists. Phylogenetic relationships were determined using the IQTree software for β -CAs from prokaryotes and protists, yellow and blue respectively. Three clades (A, B, and C) reveal regions where β -CAs from prokaryotes and protists appear to have a common ancestor.



Legend:

The conservation scale:



Variable Average Conserved

e - An exposed residue according to the neural-network algorithm.

b - A buried residue according to the neural-network algorithm.

f - A predicted functional residue (highly conserved and exposed).

s - A predicted structural residue (highly conserved and buried).

X - Insufficient data - the calculation for this site was performed on less than 10% of the sequences.

FIG 3 Sequence conservation analysis. Consurf analysis performed for the β -CA sequences in the phylogenetic tree except Clade C and protists. The conservation score is projected onto the *T. vaginalis* CA sequence. The residues strictly conserved within the Clade C sequences are indicated with violet stars (red stars used for three residues with average or low concentration).

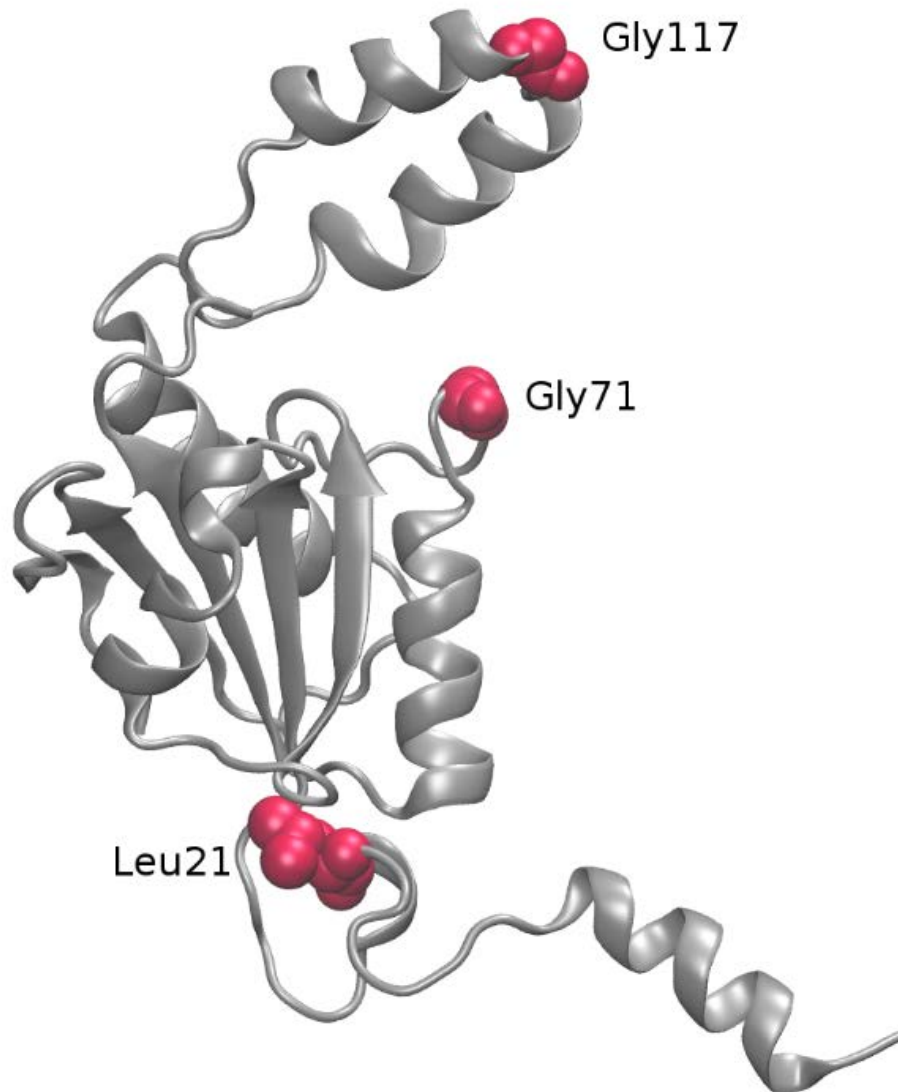


FIG 4 Evaluation of the functional importance of highly conserved residues in Clade C. A homology model of β -CA from *T. vaginalis* (A2ENQ8) (8) was used to project the conserved residues to a β -CA structure. Three conserved residues including Leu21, Gly71 and Gly117 were all located in flexible regions and are mostly exposed to solvent, indicating a non-essential structural role for these residues.