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Functional metagenomics reveals previously unrecognized diversity of antibiotic resistance genes in gulls

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Wildlife may facilitate the spread of antibiotic resistance (AR) between human-dominated habitats and the surrounding environment. Here, we use functional metagenomics to survey the diversity and genomic context of AR genes in gulls. Using this approach, we found a variety of AR genes not previously detected in gulls and wildlife, including class A and C β -lactamases as well as six tetracycline resistance gene types. An analysis of the flanking sequences indicates that most of these genes are present in *Enterobacteriaceae* and various Gram-positive bacteria. In addition to finding known gene types, we detected 31 previously undescribed AR genes. These undescribed genes include one most similar to an uncharacterized gene in *Verrucomicrobium* and another to a putative DNA repair protein in *Lactobacillus*. Overall, the study more than doubled the number of clinically relevant AR gene types known to be carried by gulls or by wildlife in general. Together with the propensity of gulls to visit human-dominated habitats, this high diversity of AR gene types suggests that gulls could facilitate the spread of AR.

Keywords: herring gulls, antibiotic resistance, *Verrucomicrobia*, *Larus argentatus*, metagenomics

INTRODUCTION

The vast majority of research on emerging antibiotic resistance (AR) has focused on AR in clinical settings. However, there is growing recognition that resistance acquired outside of hospital settings (community-acquired infections) is a serious and growing threat (Pitout et al., 2005). Antibiotics are likely important for bacterial interactions in natural communities as resistance genes originated in environmental bacteria long before the antibiotic era (Aminov and Mackie, 2007; Martinez, 2008; Davies and Davies, 2010). The collection of all resistance genes and their precursors in pathogenic and non-pathogenic microorganisms has been termed the antibiotic “resistome” (Wright, 2007).

Bacteria can readily exchange DNA via lateral transfer of mobile genetic elements (e.g., plasmids or transposons). Thus, AR genes from non-pathogenic environmental microorganisms have the potential to be transferred to pathogens infecting humans, especially when use of antibiotics imposes heavy selective pressure. Indeed, as a result of widespread antibiotic use in medicine and agriculture, multidrug resistant bacteria and resistance genes have been detected in many environments (e.g., Sayah et al., 2005; Seyfried et al., 2010). Antibiotics and AR bacteria originating from human-made AR “reservoirs” may be released into nearby groundwater, streams, rivers, and estuaries (Li et al., 2009; Martinez, 2009), potentially leading to emergence of novel mechanisms of resistance in pathogenic organisms. However, discovery of new AR mechanisms is limited because most studies focus on culturable bacteria and known genes detected via PCR. This approach has resulted in a heavily biased and incomplete understanding of the broader environmental resistome.

Wildlife may facilitate exchange of AR bacteria and genes between human-made and natural environments. AR bacteria and, more recently, AR genes have been detected in a variety of wild birds and mammals (e.g., Mallon et al., 2002; Literak et al., 2009; Marrow et al., 2009). Many species of wildlife visit human-made areas such as farms and sewage lagoons where they may acquire AR (Osterblad et al., 2001; Blanco et al., 2009). Further, some studies have discovered high abundances of AR bacteria in wildlife with little or no apparent exposure to anthropogenic sources (Williams et al., 2011).

Gulls may be a particularly important carrier of AR because of their close association with humans. During the past several decades, gull populations have increased worldwide due in large part to burgeoning human populations along coasts and the associated increase in availability of human-derived foods (Duhem et al., 2008; Schwemmer et al., 2008). Gulls carry a diverse array of potentially pathogenic bacteria including *Salmonella* spp. (Skov et al., 2008), *Campylobacter* spp. (Waldenström et al., 2007), *Listeria* spp. (Fenlon, 1985), and vero cytotoxin-producing *Escherichia coli* O157 (Wallace et al., 1997), which they likely acquire by feeding and loafing at sewage lagoons, garbage dumps, and farms (Benskin et al., 2009). Moreover, several recent studies have detected clinically relevant AR bacteria (Cizek et al., 2007; Rose et al., 2009) and AR genes (e.g., Dolejska et al., 2007; Bonnedahl et al., 2010) in gull feces.

Here, we use functional metagenomics to characterize the diversity of the gull resistome. Previous studies of AR genes in gulls and other wildlife have relied on initial cultivation of bacteria or PCR of specific, known gene variants. However, studies using functional

metagenomics have revealed a diverse array of previously undescribed AR genes in soils (Rondon et al., 2000) and humans (Diaz-Torres et al., 2006; Sommer et al., 2009). We assay: (1) the most abundant AR classes and gene types within these classes that confer resistance to four antibiotics; (2) information about the genes' genomic context (i.e., the genomic neighborhood) as revealed by the flanking sequences; and (3) microdiversity within the abundant gene types. Specifically, we screened the libraries for resistance against ampicillin, amoxicillin, penicillin, and tetracycline. These four antibiotics were chosen due their historical importance, differences in resistance mechanism, and range of target organisms. Penicillin, ampicillin, and amoxicillin are β -lactam antibiotics. In the past 20 years, bacterial strains that produced extended-spectrum β -lactamases (ESBLs) – enzymes conferring broad resistance to β -lactam antibiotics – have emerged posing a major threat to public health (Pitout and Laupland, 2008). Tetracyclines have been used for more than 40 years in clinical and agricultural settings and are broad-spectrum agents, exhibiting activity against a wide range of organisms (Chopra and Roberts, 2001). Resistance to tetracyclines is also widespread in the environment (e.g., Nascimento et al., 2003; Bryan et al., 2004).

MATERIALS AND METHODS

SAMPLING

Fecal samples were collected from four Herring Gulls in a breeding colony at the Shoals Marine Laboratory on Appledore Island, ME (42° 58' N, 70° 36' W) located approximately 10 km off the coast of mainland United States. Approximately 650 pairs of Herring Gulls nest on the island (Ellis, unpublished data). We collected fresh samples by observing individual birds until defecation occurred and immediately collecting samples from the ground using a sterile syringe. For comparison, one soil and two gull samples were also collected from the mainland (Rochester, NH, USA). Samples were immediately placed on ice. DNA extraction was performed with the FastDNA Spin for Soil kit (MP Biomedical, CA, USA) using the protocol of the manufacturer with the following changes. Cells were lysed using the MP Biomedicals FastPrep24 (setting 5.0 twice for 45 s and once for 20 s, cooling on ice in between). The precipitation step was carried out twice with 250 μ l protein precipitation solution (PPS) from the FastDNA Spin for Soil kit (MP Biomedical, CA, USA). We used 1.5 ml binding matrix and inverted for 4 min. We discarded 1 ml of the supernatant. The final pellet was resuspended in 75 μ l of the provided water.

CLONING AND SCREENING

Three micrograms of genomic DNA from gull feces was sheared to a target size of 3 kb with a Covaris S2 instrument (Covaris Inc., MA, USA). The 3-kb fragments were end-repaired with the DNA Terminator End Repair kit (Lucigen, WI, USA). The DNA was loaded on a 1% low melt agarose gel and fragments between 1 and 3 kb were extracted with Zymoclean Gel Recovery kit (Zymo Research, CA, USA). DNA was eluted in 10 μ l TE elution buffer and ligated into pSMART-HCkan vector (accession number AF532107) using the Clonesmart blunt end cloning kit (Lucigen, WI, USA). Ligation was carried out overnight at room temperature. *E. coli* 10G electrocompetent cells (Lucigen, WI, USA) were transformed by electroporation using a BTX electroporator and 1.0 mm gap cuvettes.

Electroporation conditions were 25 μ F, 200 Ω , and 1800 V. The transformation was incubated for 1 h at 37°C and, after adding 30 μ g/ml kanamycin, the transformation mixture was incubated another 2 h at 37°C. We plated 0.1 μ l of the transformation mixture on kanamycin LB plates for titer counts and 100–200 μ l on ampicillin (50 μ g/ml), amoxicillin (20 μ g/ml), penicillin (50 μ g/ml), and tetracycline (8 μ g/ml) containing plates. Screened clones were picked and grown up over night without shaking in LB + 10% glycerol. Sequencing was done using the SL1 forward primer (5'-CAGTCCAGTTACGCTGGAGTC-3') and SR2 reverse primer (5'-GGTCAGGTATGATTTAAATGGTCAGT-3'); as suggested by Lucigen) and sequenced at Beckman Coulter Genomics.

MINIMAL INHIBITORY CONCENTRATION

Antibiotic resistance clones were tested in duplicate for growth in the following concentrations of ampicillin: 60, 100, 200, 350, 500, 800, and 1000 μ g/ml. Five microliters of live culture was inoculated into 96-well plates containing 225 μ l LB Broth plus kanamycin (30 μ g/ml) and the above concentrations of ampicillin. We also included a “no-cell” control for contamination and background OD. The cultures were incubated at 37°C and growth was measured at OD₆₀₀ every 15 min for 24 h.

Tn5 TRANSPOSON “KNOCKOUT” OF AR GENES

To identify the AR gene in selected inserts, we randomly inserted a Tn5 transposon in the clones using the EZ-TN5 <TET-1> Insertion Kit (Epicentre, CA, USA). We used a molar ratio Tn5 to AR plasmid of one to minimize multiple insertion events. One microliter of a transposon reaction was electroporated (BioRad BTX) into 50 μ l of *E. Coli* 10G Electrocompetent Cells (Lucigen). Cells were plated on LB Agar containing kanamycin (30 μ g/ml) and tetracycline (10 μ g/ml). Colonies were next screened for Tn5 “knockouts” based on lack of ampicillin resistance in LB Broth plus kanamycin (30 μ g/ml), tetracycline (10 μ g/ml), and ampicillin (60 μ g/ml). A replicate control 96-plate was also inoculated with the same colony in LB Broth plus kanamycin (30 μ g/ml) and tetracycline (10 μ g/ml). Both plates were grown at 37°C (without shaking) overnight. Sequencing was done using the TN5 <TET-1> forward and reverse primers (provided in Epicentre kit).

SEQUENCE ANALYSIS

We trimmed the paired-end sequences using Phrap and analyzed the inserts using blast against both GenBank nr and the Antibiotic Resistance Genes Database (ARDB; Liu and Pop, 2009). For β -lactamases, the ARDB classification scheme generally follows other systems for organizing resistance genes (Bush and Jacoby, 2010). However, here we rely entirely on a sequence-based comparison. Using ARDB, genes are divided into the four major β -lactamase classes (A–D), then into specific ARDB types, and finally into sequence variants within each type. We detected some sequences that were identical to β -lactamases used in common cloning vectors; because we cannot determine whether these sequences are contaminants we removed them from the analysis.

Sequences from each ARDB type were assembled into contigs using Geneious (Biomatters). For selected inserts, we aligned and translated the sequences and built phylogenetic trees with Phylip (100 bootstraps) using neighbor-joining (protein distance

matrix with Jones–Taylor–Thornton correction) and maximum likelihood methods (Jones–Taylor–Thornton correction and constant rate of change among sites; Felsenstein, 2006). The phylogenetic trees were visualized with the Interactive Tree of Life (iTOL, <http://itol.embl.de/index.shtml>; Letunic and Bork, 2007). Sequences from this study have been submitted to GenBank under accession numbers JM426721–JM427844.

RESULTS

AR GENE DIVERSITY

To target AR in gulls, we sampled feces from four gulls on Appledore Island, Maine, and two gulls in Rochester, NH, USA. For comparison, we also sampled soil in Rochester without any gulls nearby (Table 1). We extracted and sheared the DNA, then cloned 3 kbp DNA fragments into *E. coli*. The number of resulting clones ranged between 1×10^5 and 4.4×10^6 per library. Based on these numbers, we estimate that up to 5% of the bacteria in gull feces are resistant to antibiotics (Table 1). We found ampicillin, amoxicillin, and penicillin resistance genes in all gulls assayed, whereas we detected tetracycline AR genes in only two out of six gulls. Although it is difficult to compare across antibiotics due to differences in potency and concentration, resistance to ampicillin was found in higher occurrence than amoxicillin and penicillin.

Gulls carried an extensive genetic diversity of both known and previously undescribed AR genes. To characterize this diversity in the metagenomic libraries, we paired-end sequenced the resistant clones and first classified the genes using ARDB (Liu and Pop, 2009; Tables 2 and 3). Overall, we found that similar genes conferred resistance to ampicillin, amoxicillin, and penicillin, including nine different class A and C β -lactamase ARDB categories. The most common category was bl1_ec, which was 100% similar to a β -lactamase in *E. coli* strain 1.1.43. Another common category detected was the extended-spectrum class A β -lactamase (ARDB category: bl2b_tem1). However, the bl2b_tem1 types were not identical to known variants in the database and on average shared only 92% sequence similarity. Combined, these two ARDB categories constituted more than 50% of the AR clones (Table 2). The last common ARDB category was the class A CTX-M type. However, these genes only shared 72% amino acid sequence similarity to CTX-M genes in *E. coli* (Table 3). A broader search in GenBank and a phylogenetic analysis revealed that the gene was 100% similar to the β -lactamase variant RAHN-2 in the Gram-negative enteric bacterium *Rahnella aquatilis* Y9602 (Ruimy et al., 2010) and therefore, was likely not a CTX-M variant (Figure 1).

Beyond the class A and class C β -lactamases, we also found ampicillin- and penicillin- resistant clones encoding for efflux

Table 1 | Metagenomic sample overview, including a list of sample IDs and description, titer of each clone library, and the number of positive (resistant) clones to each antibiotic.

Sample id	Location	Environment	Antibiotic	Total titer	+ Clones	% AR cells
10-126	Appledore	Gull feces	Ampicillin	3.7×10^6	72	3.9
10-126	Appledore	Gull feces	Amoxicillin	–	58	3.1
10-126	Appledore	Gull feces	Penicillin	–	58	3.1
10-126	Appledore	Gull feces	Tetracycline	–	4	0.2
10-131	Appledore	Gull feces	Ampicillin	9.4×10^5	8	1.7
10-131	Appledore	Gull feces	Tetracycline	–	0	0
10-145	Appledore	Gull feces	Ampicillin	4.0×10^6	20	1.0
10-145	Appledore	Gull feces	Amoxicillin	–	15	0.8
10-145	Appledore	Gull feces	Penicillin	–	20	1.0
10-145	Appledore	Gull feces	Tetracycline	–	0	0
10-186	Appledore	Gull feces	Ampicillin	5.25×10^5	5	1.9
10-186	Appledore	Gull feces	Amoxicillin	–	3	1.1
10-186	Appledore	Gull feces	Penicillin	–	2	0.8
10-186	Appledore	Gull feces	Tetracycline	–	0	0
10-240	Rochester	Gull feces	Ampicillin	9.2×10^5	19	4.1
10-240	Rochester	Gull feces	Amoxicillin	–	20	4.3
10-240	Rochester	Gull feces	Penicillin	–	23	5.0
10-240	Rochester	Gull feces	Tetracycline	–	0	0
10-241	Rochester	Gull feces	Ampicillin	1.7×10^5	N/A	N/A
10-241	Rochester	Gull feces	Amoxicillin	–	N/A	N/A
10-241	Rochester	Gull feces	Penicillin	–	N/A	N/A
10-241	Rochester	Gull feces	Tetracycline	–	4	4.7
10-245	Rochester	Soil outside gull area	Ampicillin	4.4×10^6	18	0.8
10-245	Rochester	Soil outside gull area	Amoxicillin	–	9	0.4
10-245	Rochester	Soil outside gull area	Penicillin	–	12	0.5
10-245	Rochester	Soil outside gull area	Tetracycline	–	8	0.4

Percent AR cells are calculated based on an average insert size of 2 kb and an average gull bacteria genome size of 4 Mbp.

Table 2 | The abundance of known AR genes (ARDB classification) from seven metagenomic libraries sorted by antibiotics used for selection.

Gene	Description	Organism	Genomic neighborhood	Prev. detected		Total	Amp		AX		Pen		Tet	
				Gulls ¹	Wildlife ¹		Gull N = 6	Soil N = 1	Gull N = 6	Soil N = 1	Gull N = 6	Soil N = 1	Gull N = 6	Soil N = 1
<i>acrA</i>	RND efflux pump	<i>Shigella</i>	N/A	No	No	1	1	0	0	0	0	0	0	0
B11_ampC	Class C β-lactamase	<i>Enterobacter</i>	Transcriptional regulator	No	No	2	0	0	0	0	2	0	0	0
B11_cmy2	Class C β-lactamase	<i>Enterobacteriaceae</i>	On insertion element	No	Yes	2	1	0	0	1	0	0	0	0
B11_ec	Class C β-lactamase	<i>E. coli</i>	Chromosomal	No	No	69	20	9	13	5	17	5	0	0
B11_sm	Class C β-lactamase	<i>Providencia rettgeri</i>	xre toxin system	No	No	1	0	0	0	0	1	0	0	0
B12a_1	Class A β-lactamase	<i>Bacillus</i> ⁺	Hypo. protein	No	No	1	0	0	1	0	0	0	0	0
B12a_iii	Class A β-lactamase	<i>Carnobacterium</i> ⁺	Transcriptional regulator	No	No	3	0	2	0	0	1	0	0	0
B12b_tem1	Class A β-lactamase	Many different		Yes	Yes	14	6	1	1	0	5	1	0	0
B12be_ctxm	Class A β-lactamase	<i>Rahnella</i>	Chemotaxis protein	Yes ²	Yes ²	6	5	0	1	0	0	0	0	0
B12be_oxy1	Class A β-lactamase	<i>Klebsiella</i>	Inner membrane protein	No	No	2	0	0	1	0	1	0	0	0
<i>macB</i>	RND efflux pump	<i>E. coli</i> , Gram+	Transcriptional regulator	No	No	3	0	0	0	0	1	2	0	0
<i>pbp1a</i>	Penicillin resistance	<i>Enterococcus</i> ⁺	N/A	No	No	1	1	0	0	0	0	0	0	0
<i>tetC</i>	Tet. efflux pump	<i>Clostridium</i> ⁺	ATPase	No	Yes	1	0	0	0	0	0	0	1	0
<i>tetJ</i>	Tet. efflux pump	<i>Proteus</i>	Phage lysis protein	No	No	1	0	0	0	0	0	0	1	0
<i>tetL</i>	Tet. efflux pump	<i>Enterococcus</i> ⁺	On insertion element or plasmid	Yes	Yes	1	0	0	0	0	0	0	1	0
<i>tetM</i>	Ribosomal protection	Gram+	Elongation factor G	Yes	Yes	2	0	0	0	0	0	0	2	0
<i>tetO</i>	Ribosomal protection	<i>Roseburia</i> ⁺	GTP binding protein	No	Yes	1	0	0	0	0	0	0	1	0
<i>tetW</i>	Ribosomal protection	Gram+	On insertion element	No	Yes	1	0	0	0	0	0	0	0	1
Unknown				No	No	32	14	0	8	0	3	1	4	2

N is the number of libraries screened.

¹These two columns indicate whether the AR gene type has previously been detected in either gulls or wildlife in general. This is based on the following studies: Costa et al. (2006, 2008), Dolejska et al. (2007), Bonnedahl et al. (2009, 2010), Drobni et al. (2009), Figueiredo et al. (2009), Jeters et al. (2009), Literak et al. (2009), Hernandez et al. (2010), Radhouani et al. (2010), Rybarikova et al. (2010), Silva et al. (2010), Simoes et al. (2010), Ahmad et al. (2011), Alroy and Ellis (2011).

²The *Rahnella* variant has not been observed in gulls or wildlife before.

Table 3 | Amino acid similarity of AR genes and flanking regions to sequences in ARDB and GenBank databases.

Type	ARDB			GenBank (AR gene)			GenBank (genome neighborhood)		
	Sim (%)	Organism	Best hit	Accession no.	Sim (%)	Organism	Best hit	Accession no.	Sim (%)
acrA	93	<i>Shigella flexneri</i>	acrA	ADA72786	98	N/A	N/A		
B11_ampC	85	<i>Enterobacter</i>	bla(lampC)	BAA32077	100	<i>Enterobacter</i>	Trans. regulator <i>entR</i>	ZP_05970670	90
B11_cmy2	92	<i>Citrobacter</i>	bla(CMY45)	ACV32310	95	<i>Salmonella</i>	<i>sugE</i>	YP_209396	59
B11_ec	100	<i>E. coli</i> 1.1.43	bla(lampC)	YP_002806354	100	<i>E. coli</i>	<i>b/c, frdB, frdC, frdD, sugE,</i> and <i>hyp.</i> protein	See text	90–100
B11_sm	50	<i>Providencia</i>	bla	ZP_06126521	85	<i>Providencia</i>	<i>xre</i> toxin system	ZP_06126519	97
B12a_1	100	<i>Bacillus</i>	bla	ZP_04276394	100	<i>B. anthracis</i>	Hyp. protein BA_3499	NP_845779	100
B12a_iii	49	<i>Carnobacterium</i>	bla	AAV65950	80	<i>Carnobacterium</i>	Trans. regulator	ZP_02183939	61
B12b_tem1	92								
B12be_ctxm	72	<i>Rahnella</i>	bla(RAHIN-2)	AAK83220	100	<i>Rahnella</i>	Chemotaxis protein	AAK83219	94
B12be_oxyl	85	<i>Klebsiella</i>	bla	CAQ76676	85	<i>Klebsiella</i>	Inner membrane protein	YP_001335817	
macB	37		ABC transporter	YP_817928 or ZP_071822747	95		Trans. regulator	ZP_04782719 or ZP_07165374	70–90
pbp1a	45	<i>Enterococcus</i>	Glycosyl transferase	EFM68814	89	N/A	N/A	N/A	N/A
tetC	100	Plasmid	IncN	NP_511233	100	<i>Clostridium</i>	ATPase	ZP_02633149	98
tetJ	54	<i>Proteus</i>	tetJ	ZP_03841720	100	<i>Proteus</i>	phage lysis protein	YP_002152118	100
tetL	96	<i>Enterococcus</i>	Tn916	YP_133685	100	<i>Enterococcus</i>	Hyp. protein	ZP_05563488	100
tetM	100	<i>Streptococcus</i>	tetM	ADM91220	95	<i>Enterococcus</i>	Elongation factor G	EFM68952	95
tetO	100	<i>Streptococcus</i>	tetO	ACT76131	96	<i>Roseburia</i>	GTP binding protein	CBL13258	93
tetW	92	<i>Lawsonia</i>	tetW	YP_594556	96	<i>Lactobacillus</i>	Transposase	YP_003601920	100
unk1	N/A	<i>Lactobacillus</i>	Hydrolase	EFG55459	48	<i>Enterococcus</i>	PIIT domain protein	ZP_03985770	84
unk2	N/A	<i>Burkholderia</i>	Flagellar asso. protein	ZP_02357438	53	<i>Nakamurella</i>	UDP-glucosyltransferase	YP_003204087	71
unk3	N/A	<i>Methylobacterium</i>	Hypothetical protein	YP_001691243	79	<i>Rhizobium</i>	Hypothetical protein	YP_471752	60
unk4	N/A	<i>Enterococcus</i>	Dihydroorotate dehydrogenase	ZP_06697492	69	<i>Mycoplasma</i>	Dihydroorotate dehydrogenase	NP_758176	68
unk5	N/A	<i>Pseudomonas</i>	Hypothetical protein	YP_418084	52	<i>Pseudomonas</i>	Hypothetical protein	YP_418084	52
unk6	N/A	<i>E. coli</i>	Methionyl-tRNA synthetase	ZP_07099092	93	<i>E. coli</i>	Methionyl-tRNA synthetase	ZP_07099092	100
unk7	N/A	<i>Enterococcus</i>	Hypothetical protein	YP_398703	79	<i>Enterococcus</i>	Hypothetical protein	YP_398703	79
unk8	N/A	<i>Enterococcus</i>	Hydrolase	EFM78292	56	<i>Enterococcus</i>	Hydrolase	EFM78292	56
unk9	N/A	<i>Citrobacter</i>	ATP synthase subunit	YP_003367440	99	<i>Citrobacter</i>	ATP synthase subunit	YP_003367440	99
unk10	N/A	<i>E. coli</i>	Hpt domain protein	ZP_07146767	97	<i>E. coli</i>	Hpt domain protein	ZP_07146767	100
unk11	N/A	<i>Enterococcus</i>	NOL1/NOP2/sun family protein	ZP_05646561	85	<i>Enterococcus</i>	NOL1/NOP2/sun family protein	ZP_05646561	83

(Continued)

Table 3 | Continued

Type	ARDB			GenBank (AR gene)			GenBank (genome neighborhood)		
	Sim (%)	Organism	Best hit	Accession no.	Sim (%)	Organism	Best hit	Accession no.	Sim (%)
unk12	N/A	<i>Bacillus</i>	Uracil-DNA glycosylase	YP_001377098	74	<i>Burkholderia</i>	Short-chain dehydrogenase	YP_002909251	95
unk13	N/A	<i>E. coli</i>	Multidrug transporter	ZP_06938716	93	<i>E. coli</i>	Multidrug transporter	YP_002385928	91
unk14	N/A	<i>Streptococcus</i>	Integrase	ZP_04061436	91	<i>Enterobacter</i>	AraC family transcriptional regulator	YP_001175810	82
unk15	N/A	<i>Verrucomicrobium</i>	Hypothetical protein	ZP_02929363	84	<i>Gemmata</i>	Protease	ZP_02733734	45
unk16	N/A	<i>Arcobacter</i>	Auxin efflux protein	YP_001489985	92	N/A	N/A	N/A	N/A
unk17	N/A	<i>P. putida</i>	Alcohol dehydrogenase	YP_001668264			panetheine-phosphate adenylyltransferase	ZP_07057662	
unk18	N/A	<i>Photobacterium</i>	Hypothetical gene	ZP_06155446	93	<i>Pseudomonas</i>	TirH transporter	YP_791383	80
unk19	N/A	<i>E. coli</i>	Hypothetical gene	CAA74339	70	<i>E. coli</i>	IS1 transposase B	NP_414562	91
unk20	N/A	<i>Bacillus</i>	β -Lactamase precursor	ZP_06873125	47	<i>Bacillus</i>	Hypothetical protein	ZP_03104389	60
unk21	N/A	<i>Proteus</i>	Chondroitin ABC lyase precursor	ZP_03840872	93	<i>Proteus</i>	Chondroitin ABC lyase precursor	YP_002151847	87
unk22	N/A	<i>Bacteroides</i>	Tyrosine type site-specific recombinase	ZP_06094951	98	<i>Bacteroides</i>	Tyrosine type site-specific recombinase	YP_100741	88
unk23	N/A	<i>Proteus</i>	Colicin V production protein	YP_002151514	85	<i>Proteus</i>	Hypothetical protein	YP_002151515	82
unk24	N/A	<i>Providencia</i>	TPR repeat protein	ZP_05971549	62	<i>Proteus</i>	Hypothetical protein	ZP_03802426	85
unk25	N/A	<i>Proteus</i>	Iron ABC transporter	YP_002152660	93	<i>Proteus</i>	Iron ABC transporter	YP_002152659	94
unk26	N/A	<i>Lactobacillus</i>	Transposase	ZP_05866387	94	<i>Faecalibacterium</i>	Hypothetical protein	ZP_02091680	96
unk27	N/A	<i>Pantoea</i>	Phosphoglycerate mutase	ZP_07378382	94	<i>Yersinia</i>	Hypothetical protein	ZP_04630452	88
unk28	N/A	<i>Streptomyces</i>	Hypothetical protein	NP_822969	74	N/A	N/A	N/A	N/A
unk29	N/A	<i>Bacillus</i>	Histidine kinase	YP_175245	100	N/A	N/A	N/A	N/A
unk30	N/A	<i>Clostridium</i>	D-3-phosphoglycerate transferase	ZP_02639608	100	<i>Clostridium</i>	D-3-phosphoglycerate transferase	ZP_02639608	100
unk31	N/A	<i>Lactobacillus</i>	DNA repair gene	ZP_05752538	92	<i>Lactobacillus</i>	Hypothetical protein	ZP_03963367	68

pumps (*acrA* and *macB* type) and a membrane carboxypeptidase (ARDB category: *pbp1a*). Finally, we observed a high proportion of clones (38%) that had no clear match in ARDB.

The tetracycline resistant genes detected belonged to several ARDB categories including *tetC*, *tetJ*, and *tetL* efflux pumps and *tetM*, *tetO*, and *tetW* ribosomal modification proteins. These genes were generally very similar to known genes (Table 3) and likely do not constitute new variants.

For comparison, we also analyzed the genetic diversity of AR in a nearby soil sample. Keeping in mind that we only analyzed one soil sample, we found a lower frequency of AR genes in the soil sample compared to the gulls. With the exception of *tetW*, all ARDB categories observed in soil were also detected in gulls. Furthermore, we did not detect the RAHN-2/CTX-M category in soil. Thus, it appears that many AR genes in gulls are also found in soil (and vice-versa), with perhaps some exceptions.

GENOMIC NEIGHBORHOOD

Antibiotic resistance genes are known to move around between hosts via lateral gene transfer, thus the AR gene sequence itself is not a good indicator of the organism with the gene. Therefore, we analyzed the flanking sequences around the AR genes (Figure 2; Tables 2 and 3). Data regarding the genomic neighborhood can give information about the host organism and/or whether the gene was located on a mobile genetic element. Our attempts to assemble clones belonging to the same ARDB category yielded good contigs, indicating that the detected AR genes shared flanking sequences. An example of that was the common class C β -lactamase belonging to the ARDB category *bl1_ec*. All clones associated with this category were located on a genomic fragment syntenous with *E. coli* strain K-12 substr. DH10B and several other *E. coli* strains (Figure 2A). Similar to DH10B, the gene was flanked by fumarate reductase subunits *frdB*, *frdC*, and *frdD*, an outer membrane lipoprotein (*Blc*), a quaternary ammonium compound-resistance protein (*sugE*), and the toxin entericidin (Durfee et al., 2008). Overall, this suggests that the most common class C β -lactamase found in the gulls was located on a genomic fragment similar to that found in *E. coli*. Similarly, the β -lactamase variant RAHN-2 also assembled into one contig which suggested a fragment syntenous with the genome from *R. aquatilis* Y9602 (Figure 2B). This contig including a chemotaxis gene and a hypothetical protein. This supports the hypothesis that this variant is indeed located in a *Rahnella* genome.

In contrast to the *bl1_ec* and RAHN-2 variants, clones matching the TEM class A β -lactamases (*bl2b_tem1*) were found in a range of genetic neighborhoods. The genes flanking *bl2b_tem1* sequences were in some cases most closely related to Gram-positive bacteria (e.g., *Lactobacillus*, *Enterococcus*, and *Clostridium*) and in other cases, to Gram-negative bacteria (e.g., *E. coli*, *Shigella*, and *Salmonella*).

There were two major types of tetracycline resistance genes in our samples – efflux pumps (*tetC*, *tetJ*, and *tetL*) and ribosomal modification proteins (*tetM*, *tetO*, and *tetW*; Table 2). With the exception of *tetJ*, these were all associated with Gram-positive bacteria including *Bacillus*, *Enterococcus*, and *Clostridium*. Several of them were located on what appeared to be mobile elements including plasmids and transposons.

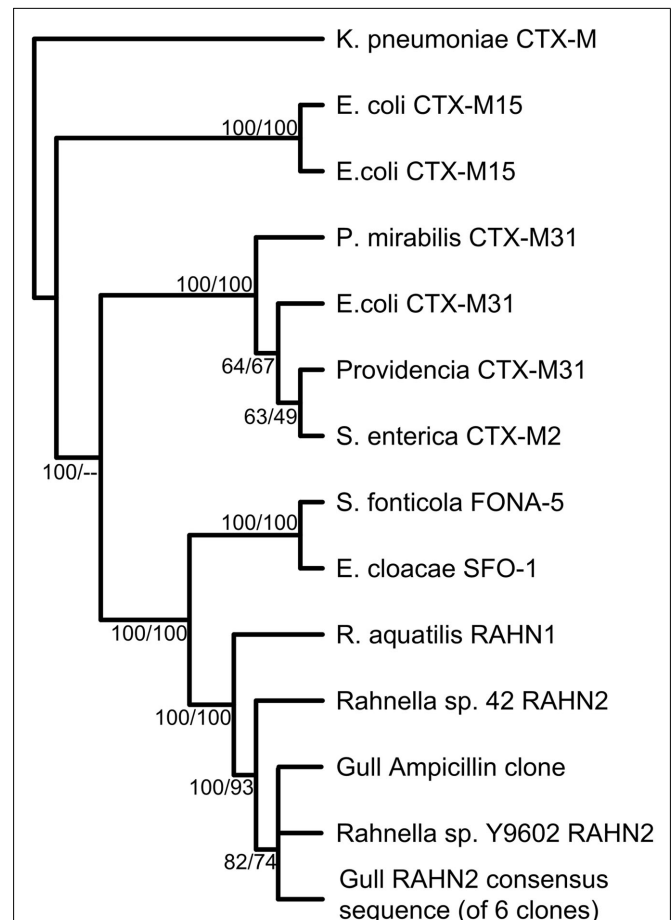


FIGURE 1 | Phylogenetic tree of the *bl2be_ctxm* type class A β -lactamase gene from metagenomic clones. The phylogenetic tree is a majority rule consensus tree based on protein similarity using neighbor-joining. Bootstrap values (total 100) are calculated with neighbor-joining and maximum likelihood methods. *Klebsiella pneumoniae* CTX-M is the outgroup.

AR GENE MICRODIVERSITY

In addition to detecting multiple classes of AR genes, we found several DNA polymorphisms in the *bl1_ec* type. To further explore this result, we aligned the sequences matching this ARDB type and built a phylogenetic tree (Figure 3). All gull *bl1_ec* sequences grouped with *E. coli* 1.1.43 with high bootstrap support. In contrast, sequences retrieved from the soil samples formed three separate clades, with only one soil clone clustering with the gull variants. Instead, the one clade of soil AR sequences were 100% similar to *E. coli* H120, whereas the other genes branched away from known *E. coli* genes. Overall, this indicates that the *bl1_ec* diversity is related to its environmental origin. We also analyzed the sequence variation among clones from the RAHN-2 β -lactamase genes but only observed one variant.

CHARACTERIZATION OF UNDESCRIBED AR GENES

In addition to the known AR gene categories in ARDB, we detected 31 gene types not previously described (Table 3). The

novel AR genes were associated with a wide range of phylogenetic lineages including *Pseudomonas*, *Bacteroides*, Gram-positives, *Verrucomicrobia*, and different genera affiliated with *Enterobacteriaceae*. To further characterize these, we first measured the minimum inhibitory concentration (MIC) for ampicillin. All ampicillin/penicillin/amoxicillin resistant clones had an MIC above 1000 µg/ml ampicillin and thus clearly conferred resistance to this antibiotic.

Two of these clones assembled to a 2800-bp DNA fragment (unk15 in **Table 3**). One end of the fragment shared 84% amino acid similarity to the hypothetical protein VspiD_21980 in *Verrucomicrobium spinosum* from the phylum *Verrucomicrobia*. The other part of the fragment was 45% similar to a C-terminal protease in *Gemmata obscuriglobus* from the phylum *Planctomycetes*.

We next identified the genomic region in the clone that encoded for AR by randomly inserting a Tn5 transposon followed by testing for ampicillin resistance and DNA sequencing of the clones. The Tn5 “knockout” revealed that insertions in both genes interrupted the resistance whereas several insertions between the two genes had no effect. This represents a completely novel AR gene system that confers resistance in *E. coli*, but at present, we do not know the specific mechanism. To the best of our knowledge, this is the first study to identify an AR gene in either the phylum *Verrucomicrobia* or *Planctomycetes*.

A second previously uncharacterized AR gene (unk1) shared 85% amino acid similarity to a protein in the Gram-positive *Carnobacterium* sp. AT7 that was annotated as a putative DNA repair protein. A Tn5 insertion in this gene interrupted the resistance. Next to this putative AR gene was a gene 85% similar to *piIT* in another Gram-positive lineage, *Enterococcus faecalis*. Another previous unrecognized AR gene (unk31) was 92% similar to a DNA repair gene of the ImpB/MucB/SamB family in *Lactobacillus coryniformis* subsp. *torquens* KCTC 3535. Multiple Tn5 insertions in this gene also interrupted the resistance. None of these DNA repair genes have previously been shown to confer AR, so the exact mechanism of resistance is unclear.

DISCUSSION

Using a functional metagenomics approach, we detected an extensive diversity of AR genes in gulls, including both well-known and undescribed AR genes. Less than half of the gene types detected have been reported previously in gulls (**Table 2**). We also detected

several completely unknown AR genes including one associated with the lesser-known phyla *Verrucomicrobia* and *Planctomycetes*. These results highlight the strength of the functional metagenomic approach: the ability to reveal the most abundant genes conferring resistance to a particular antibiotic, without requiring prior knowledge of the gene classes of interest. In contrast, most studies of antimicrobial resistance have used PCR to characterize specific AR genes, often known from clinically relevant bacterial isolates (such as, *E. coli*, *Salmonella*, *Enterococcus* spp. – see references in **Table 2**). Our results suggest that these previous studies, while useful, may impart a heavily biased view of the AR genes carried by wild animals.

The functional metagenomic approach also has several advantages over a straight sequencing-based metagenomic analysis of a community. By including a functional screening, we can identify previously unknown AR genes that we would not have been able to identify by sequence homology searches (e.g., Sommer et al., 2009). Further, most metagenomic analyses only target the most abundant lineages, whereas the functional metagenomic approach allows for screening even rare members in the community.

Of course, there are limitations to the method used here. The approach is unlikely to detect mechanisms that require large (>3 kb) gene cassettes or that confer resistance by modifying regulation of genes in another genomic region (Viveiros et al., 2007). Further, it will only find AR genes that confer resistance in *E. coli*. Nonetheless, 42% of the gene types we detected originated from Gram-positive hosts (e.g., *Enterococcus* and *Clostridium*) as well as lesser-studied phyla like *Verrucomicrobia* and *Bacteroidetes*. This indicates that this bias may not be as dramatic as would be expected. Further, the functional assay yields the additional useful information that the genes are expressed in *E. coli* and therefore have the potential to be transferred and functional in human pathogens.

Wild animals are probably natural reservoirs of AR bacteria and genes, carrying them as part of their commensal flora (Wright, 2010). Alternatively, animals may acquire AR genes through exposure to human activities, including agriculture and animal husbandry (Osterblad et al., 2001). Without a much broader understanding of natural AR gene diversity, we cannot determine the source of AR genes in gulls. However, the gulls sampled in this study carried a variety of AR genes that are commonly found in bacteria isolated from humans and domestic animals. This

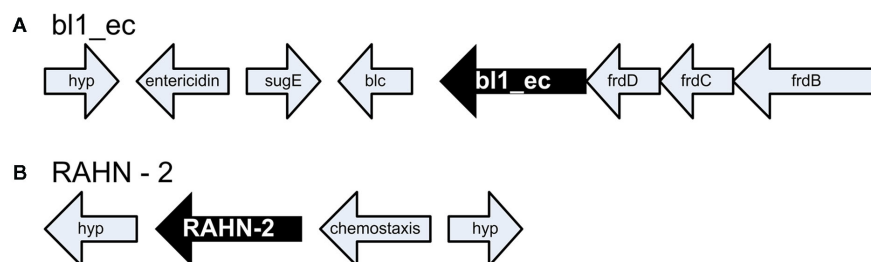
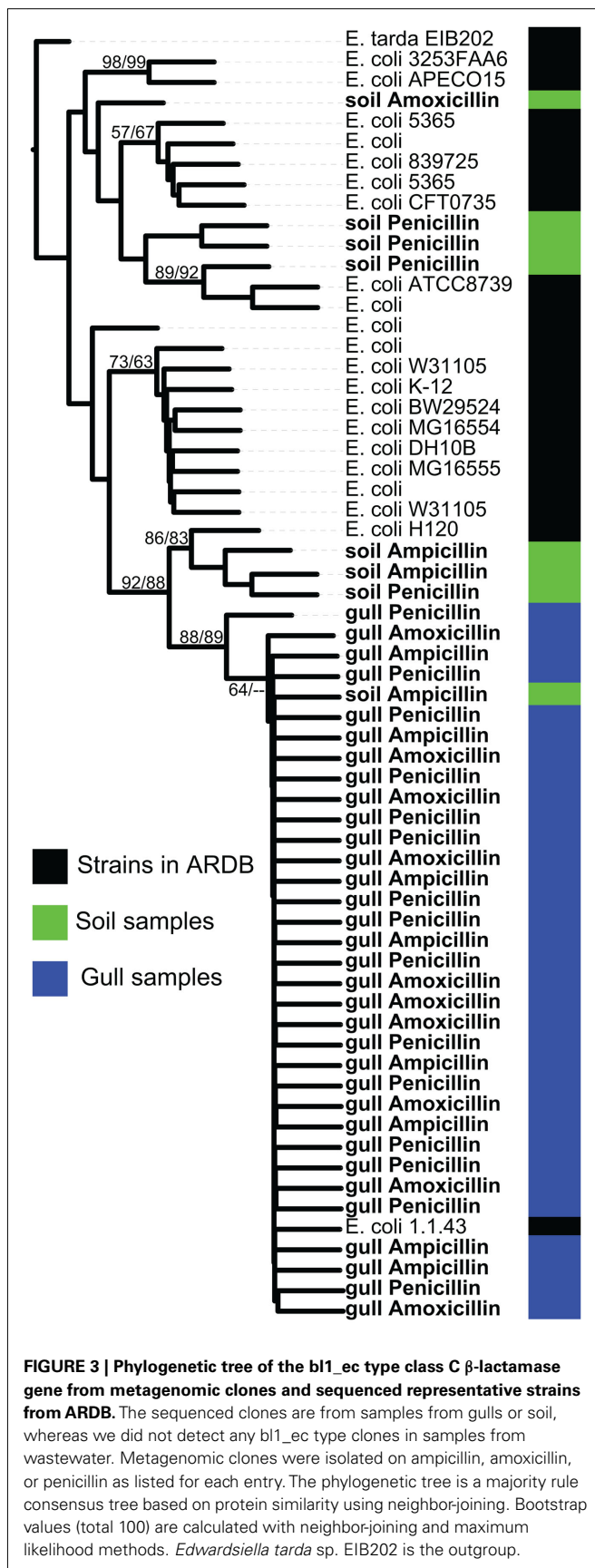


FIGURE 2 | Genome neighborhood of antibiotic resistance genes. (A) Genomic configuration of *bl1_ec*. *frdBCD* encode fumarate reductase subunits, *Blc* encodes an outer membrane lipoprotein, *sugE* encodes a

quaternary ammonium compound-resistance protein and *hyp*. encodes a hypothetical protein (B) *bl2be_ctxm*. *Hyp*. encodes two hypothetical proteins adjacent to the β -lactamase *RAHN-2* in *Rahnella*.



result, similar to that of other recent studies of gulls and other wildlife, suggests that AR genes arising from the use of antibiotics in humans and animals may have spread to wildlife. The genes may then be further disseminated by wild animals, particular those that migrate long distances such as birds (Sjolund et al., 2008; Drobni et al., 2009).

Studies of gull ecology conducted at Appledore Island, Maine (where most of our samples were taken) indicate that gulls not only carry AR genes, but also have the potential to be a vector of the genes between human-dominated habitats and the environment. Appledore-banded gulls have been found at a variety of human habitats across New England and as far away as Florida (Ellis, unpublished data). They have been observed loafing at landfills and wastewater treatment plants and share strains of *E. coli* in common with wastewater from these sites (Nelson et al., 2008). Garbage brought back from the mainland is also a large part of the diet of gulls on Appledore Island (Ellis et al., 2005). Finally, the birds are frequently observed at public beaches in coastal New Hampshire and Maine during the summer, where they come into close contact with humans. Thus, the combination of the traveling patterns of gulls and the extensive diversity of AR genes that they carry suggest that gulls may be important vectors for spreading AR genes in areas where humans live.

The most common gene detected in our study was bl1_{ec}, a class C β-lactamase. The microdiversity within this gene type provides further evidence that gulls could move AR genes from human habitats to the environment. The gull variants were highly similar (some sharing 100% aa identity) to that of *E. coli* 1.1.43, an isolate from a human gastrointestinal tract. In contrast, while the soil sample contained the gull variant, it also contained sequence diversity that did not match any known clinical isolates. Hence, gull AR gene diversity does not appear to simply reflect the diversity found in soils (and therefore, perhaps, the broader environment), although further work is needed to confirm this hypothesis.

Class A β-lactamases were also common in our samples. In a recent study of AR in Alaskan soil, class B β-lactamases were most common (Allen et al., 2009). We did not detect any class B types from our samples, however, indicating that the relative abundance of β-lactamase classes may vary by environment or location.

Only two of the ARDB β-lactamase gene types that we identified have been reported previously in gulls (Table 2). Notably, one of these genes was classified as a CTX-M-like ESBL. ESBL-producing organisms are a major public health concern in clinics and communities (Pitout and Laupland, 2008). However, the variant was also 100% similar to that previously found in *R. aquatilis* (*Enterobacteriaceae*), and the flanking sequences also suggest a *Rahnella* host. *Rahnella* has been implicated in several instances of infectious disease in humans, causing bacteremia (Chang et al., 1999; Tash, 2005). *R. aquatilis* has also been found in the guts of insects (e.g., Morales-Jimenez et al., 2009) and a wild fish (Skrodenyte-Arbaciauskiene et al., 2008); however, as far as we know, this is the first report of the genus in a wild bird.

Although we generally detected few tetracycline resistance genes in our samples, we identified five known tetracycline resistance gene types in gulls and a sixth type in soil. *tetL* and *tetM* have been reported before in a few studies of gulls and other wildlife (Table 1). Notably we did not detect *tetA* and *tetB*, which have

been widely reported in other wildlife (e.g., Silva et al., 2010; Alroy and Ellis, 2011). The gene *tetW* has only been reported in one study of wild animals (Jeters et al., 2009). *tetC* has also only been reported in one other study of wildlife; the gene was found in small rodents trapped at swine farms, suggesting that the presence of the gene may have resulted from agricultural use of antibiotics. Interestingly, several other studies were unable to detect *tetC* in wildlife using PCR (e.g., Rybarikova et al., 2010; Ahmad et al., 2011). One other study of wild animals detected *tetO* in houseflies and German cockroaches at swine farms (Ahmad et al., 2011). Thus, gulls from Appledore Island may have acquired some *tet* genes from agricultural sources.

In addition to these well-known AR genes, both gulls and soil also harbor a variety of previously unknown AR genes. Given that the genes confer resistance in *E. coli*, they clearly have the potential to be functional in human pathogens. Some of these genes may have originally derived from human bacteria and have yet to be detected in clinical settings. Indeed, Sommer et al. (2009) detected a variety of novel AR genes in healthy humans using the

same method. Alternatively, the genes may be intrinsic to the birds' commensal flora or more broadly, to the environment. In this case, the mixing of human and gull fecal bacteria could result in novel combinations and mechanisms of AR in human beings.

In conclusion, by using a functional metagenomics approach, this study more than doubled the number of clinically relevant AR gene types known to be carried by gulls or by wildlife in general. Together with the propensity of gulls to visit human-dominated habitats, this high diversity of AR gene types suggests that gulls can facilitate the spread of AR. Further, the large number of novel genes that we identified suggests that gulls also have the potential to introduce new mechanisms of AR back to the human microflora.

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