Characterization of Acinetobacter baumannii Copper Resistance Reveals a Role in Virulence

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Acinetobacter baumannii is often highly drug-resistant and causes severe infections in compromised patients. These infections can be life threatening due to limited treatment options. Copper is inherently antimicrobial and increasing evidence indicates that copper containing formulations may serve as non-traditional therapeutics against multidrug-resistant bacteria. We previously reported that A. baumannii is sensitive to high concentrations of copper. To understand A. baumannii copper resistance at the molecular level, herein we identified putative copper resistance components and characterized 21 strains bearing mutations in these genes. Eight of the strains displayed a copper sensitive phenotype (pcoA, pcoB, copB, copA/cueO, copR/cusR, copS/cusS, copC, copD); the putative functions of these proteins include copper transport, oxidation, sequestration, and regulation. Importantly, many of these mutant strains still showed increased sensitivity to copper while in a biofilm. Inductively coupled plasma mass spectrometry revealed that many of these strains had defects in copper mobilization, as the mutant strains accumulated more intracellular copper than the wild-type strain. Given the crucial antimicrobial role of copper-mediated killing employed by the immune system, virulence of these mutant strains was investigated in Galleria mellonella; many of the mutant strains were attenuated. Finally, the cusR and copD strains were also investigated in the murine pneumonia model; both were found to be important for full virulence. Thus, copper possesses antimicrobial activity against multidrug-resistant A. baumannii, and copper sensitivity is further increased when copper homeostasis mechanisms are interrupted. Importantly, these proteins are crucial for full virulence of A. baumannii and may represent novel drug targets.

Keywords: Acinetobacter baumannii, metal, copper, pathogenesis, Galleria

INTRODUCTION

Acinetobacter baumannii is responsible for a significant proportion of nosocomial infections worldwide, and an even greater number of ICU-acquired infections; co-morbidities are an important risk factor for A. baumannii infection (Vincent et al., 2009; Lob et al., 2016; Wong et al., 2017; Du et al., 2019). Associated types of infections are diverse, and include pneumonia, urinary tract infections, bacteremia, skin and soft tissue infections, osteomyelitis, and meningitis...
Extensive drug-resistance is common among *A. baumannii* strains, and resistance limits treatment options and leads to higher morbidity and mortality (Antunes et al., 2014; Giammanco et al., 2017; Du et al., 2019). Indeed, multidrug-resistant *A. baumannii* has been named a “serious” threat by the CDC since 2013, and in 2017 carbapenem-resistant *A. baumannii* topped the World Health Organization’s Priority Pathogen’s List as a “Level 1: Critical priority” pathogen (Centers for Disease Control and Prevention, 2013; World Health Organization, 2017). Clearly, new therapeutic options are desperately needed to treat multidrug-resistant *A. baumannii* infections.

Antibiotics target essential functions for bacterial growth and/or survival. Metal homeostasis is a vital process that provides an extensive list of potential new antimicrobial targets. Copper is required for cellular function, e.g., for redox balance and as an enzyme cofactor. However, copper ions become toxic at high concentrations; thus, it is important that intracellular copper levels be tightly controlled. Copper ions cause damage by participating in Fenton-like chemistry to produce hydroxyl radicals that react with and damage essential biomolecules (Liochev and Fridovich, 2002) and also by displacing iron from crucial iron-sulfur cluster proteins (Macomber and Imlay, 2002). Studies in *Escherichia coli* and *Salmonella* spp. have shown that when bacteria are placed on copper surfaces, outer membrane integrity is compromised, hydroxyl radicals are produced, respiration is inhibited, and DNA is degraded (Warnes et al., 2012). Fenton-like chemistry-based killing of pathogens also occurs in the host via host-generated reactive oxygen species. Because of the need for new therapeutics to treat antibiotic resistant pathogens, research into the use of copper as an antimicrobial has lately increased. For example, the use of copper-containing surfaces in hospitals has been shown to greatly reduce environmental contamination with nosocomial pathogens and to reduce rates of health-care-acquired infections (Salgado et al., 2013; Sifri et al., 2016; von Dessauer et al., 2016). Additionally, copper-containing wound dressings are in development to aid in healing of infected wounds (Borkow et al., 2010; Ahire et al., 2016).

The damaging effects of copper have been harnessed by the host immune system. Indeed, phagocytic immune cells employ a copper burst within the phagosome to kill pathogens (Sheldon and Skaar, 2019). Concentrations of copper upwards of 0.5 mM have been measured in macrophage phagosomes (Wagner et al., 2005). Additionally, host mobilization of copper also occurs in response to infection: increased concentrations of copper have been measured in serum and in wound exudate (Milanino et al., 1993; Jones et al., 2001). Consequently, pathogens with mutations in crucial copper resistance genes have been found to have impaired intracellular survival, colonization, and/or virulence (Djokot et al., 2015). Furthermore, it was recently shown that a copper sensitive mutant strain of *A. baumannii* demonstrated reduced colonization of the respiratory tract of mice (Alquethamy et al., 2019).

In bacterial species where copper homeostasis has been well-characterized, a variety of proteins are utilized to facilitate copper homeostasis. Though this process remains poorly understood in *A. baumannii*, recent studies have identified many putative copper resistance gene homologs in *A. baumannii* clinical isolates (Hernandez-Montes et al., 2012; Williams et al., 2016; Alquethamy et al., 2019). For example, in the model clinical multidrug-resistant isolate, AB5075, copper resistance genes exist in four distinct chromosomal regions named A–D (Figure 1) (Williams et al., 2016). These regions contain genes predicted to encode homologs of the most important copper resistance proteins of bacteria, e.g., copper ATPases, copper oxidases, copper transporters, copper chaperones, and regulatory proteins (Williams et al., 2016).

Region A includes genes for a large RND family efflux pump, *czcCBAD*, and a hypothetical protein ABUW_0265, which is located at the start of the operon. A similar type of pump, CusCFBA, has been shown to contribute to copper resistance in *E. coli* (Franke et al., 2003). While the genes in this operon are annotated as copper-related in AB5075, expression does not increase in response to copper exposure (Williams et al., 2016) and it has been suggested that *A. baumannii* in fact does not encode *cus* homologs (Alquethamy et al., 2019), thus the role of this operon in copper resistance is very tentative.

Region B encodes four genes, including homologs of two known copper resistance proteins. CueR, CopA1, and the hypothetical protein ABUW_2705 are encoded as an operon; divergently transcribed from that operon is a putative copper chaperone, ABUW_2708, recently named CopZ (Alquethamy et al., 2019). CueR has been well-studied in *E. coli* and is a cytoplasmic transcription factor that binds copper ions with exquisite sensitivity (Changela et al., 2003). Upon copper binding, CueR positively regulates transcription of copper resistance genes. In *E. coli*, CueR is known to regulate the expression of copper ATPase *CopA* and the copper oxidase *CueO* (Outten et al., 2000; Yamamoto and Ishihama, 2005). The CueR regulon in *A. baumannii* is not yet characterized, however, a putative CueR binding site has been identified in the intergenic space of region B, suggesting that all of the genes in region B may be regulated by CueR (Alquethamy et al., 2019). Appropriately, the genes of region B are all upregulated in response to copper in a similar manner (Williams et al., 2016). A CopA1 homolog, also encoded in region B, is a putative copper ATPase, which is another important and well-characterized copper resistance protein. Copper ATPases use energy from ATP to move copper ions into the periplasmic space from the cytoplasmic membrane and are known copper resistance proteins. CueR, CopA1, and the regulatory protein *CueO* have been well-studied in *E. coli* (Franke et al., 2003). While the genes in this operon are annotated as copper-related in AB5075, expression does not increase in response to copper exposure (Williams et al., 2016) and it has been suggested that *A. baumannii* in fact does not encode *cus* homologs (Alquethamy et al., 2019), thus the role of this operon in copper resistance is very tentative.

Region C encodes a single operon of five genes, ABUW_3226 – 3230. Two of the genes, ABUW_3227 and ABUW_3228, are annotated as *pcaB* and *pcoA*, respectively. In *E. coli*, the *pco* system is plasmid-based; while the *Pco* proteins seem to be homologous to known chromosomally encoded copper resistance factors, their functions are less well-studied and understood (Lee et al., 2002). In AB5075, *PcoB* is homologous to *CpcB*, an outer membrane protein that may function...
FIGURE 1 | Organization of regions A–D in the ABS0575 chromosome. We previously identified 22 putative copper-related genes in the genome of ABS0575 (Williams et al., 2016). These genes are located in four chromosomal regions named A–D. The relative location of each region on the ABS0575 chromosome is shown on the left (not to scale), and the genes and regions are depicted to scale relative to one another. Each operon is colored uniquely, and the same colors are maintained in Figure 9. Gene annotations are shown when available, otherwise ORF numbers are used.

as a copper transporter (Cha and Cooksey, 1993). PcoA is homologous to CueO, a periplasmic copper oxidase that detoxifies by oxidizing copper ions to their less damaging form (Grass and Rensing, 2001). Surprisingly, expression of the genes in region C does not change in response to copper exposure (Williams et al., 2016), however that does not preclude the possibility that these genes are indeed involved in copper resistance.

Region D is the largest of the identified regions in ABS0575; it contains a total of eight genes arranged as two pairs of divergently transcribed genes/operons. Among these eight genes are seven homologs of characterized copper resistance proteins: CopB, CueO, CusR, CusS, CopA2, CopC, and CopD. The functions of homologs of some of these copper-related proteins have been previously elucidated in other bacteria. CopB and CopD are membrane proteins and putative copper transporters, though not yet well-characterized (Cha and Cooksey, 1993). CueO is a periplasmic copper oxidase, which detoxifies by oxidizing cuprous copper ions (Cu$^{1+}$) to their less damaging cupric (Cu$^{2+}$) form (Grass and Rensing, 2001). CopA2 is a second copper ATPase (the other being CopA1 of region B); previous work did not reveal a role for copA2 in copper resistance in ABS0575 (Alquethamy et al., 2019). CopC is a periplasmic copper chaperone protein that likely contributes to copper resistance by binding copper ions as a means to prevent reactivity and/or by shuttling ions between other copper resistance proteins (Puig et al., 2002; Arnesano et al., 2003; Padilla-Benavides et al., 2014). CusR and CusS form a two-component system that senses periplasmic copper ions and activates the CusR regulon in response (Gudipaty et al., 2012). In E. coli, CusR is known to directly regulate expression of the cusRS operon and the divergently transcribed cusCFBA operon (Munson et al., 2000; Yamamoto and Ishihama, 2005; Gudipaty et al., 2012). It is not yet known which genes comprise the CusR regulon in A. baumannii, however, putative CusR binding sites were identified in the intergenic regions upstream of the four operons in region D, suggesting that all of region D may be regulated by CusR (Alquethamy et al., 2019).

Notably, not all clinical isolates of A. baumannii carry all of the copper-related genes found in ABS0575. Based on previous analyses of two panels of clinical isolates, as well as the common lab strain ATCC 17978, region A–C genes are encoded by all strains. Conversely, region D genes are found in about 40% of strains (5/12 strains) (Williams et al., 2016; Hassan et al., 2017). Furthermore, the presence of region D genes is linked to increased copper resistance. Indeed, isolates show marked differences in copper sensitivity that directly correlates with the presence of the region D genes; clinical isolates lacking region D demonstrate significantly less copper resistance than isolates that encode the region D genes (Williams et al., 2016). Moreover, expression of the genes in region D dramatically increases in response to copper exposure (Williams et al., 2016). These results strongly suggest that the region D genes are crucial for extensive copper homeostasis capability in A. baumannii.

Given the growing interest in targeting copper homeostasis as a novel antimicrobial strategy and the lack of understanding of which factors contribute to copper resistance in A. baumannii, we sought to identify genes that were important for copper resistance in this important pathogen. Herein, we describe a detailed analysis of 21 mutant strains, each bearing insertions in the genes carried in regions A–D (Figure 1). Marked copper
sensitivity of a subset of the mutant strains as well as defects in virulence are described.

**MATERIALS AND METHODS**

**Bacterial Strains and Growth Conditions**

Bacterial strains are listed in Table 1. Strains were routinely streaked from freezer stocks and were grown overnight at 37°C on lysogeny broth (LB) 1.5% agar plates (MoBio, Carlsbad, CA, United States). Liquid cultures of *A. baumannii* were grown at 37°C shaking at 190 rpm in LB or in M9 minimal medium (Sambrook and Russell, 2001) supplemented with 0.1% casamino acids (Difco, Franklin Lakes, NJ, United States); the M9 medium was always supplemented with amino acids, but for simplicity is referred to as 'M9 medium' throughout the manuscript. Importantly, overnight cultures were consistently started from a single opaque colony, as phase variation has been shown to affect growth and virulence (Tipton et al., 2015). Overnight growth was restricted to a period of 16–19 h. *A. baumannii* strains freezer stocks were maintained at −80°C in medium supplemented with 40% glycerol.

**AB5075 Mutant Strains**

Transposon mutant strains were purchased from the comprehensive, ordered transposon library engineered by Dr. Colin Manoil's group at the University of Washington (Gallagher et al., 2015). In general, we chose strains with transposon insertions near the 5' end of the gene; most have insertions in the first third of the ORF (Table 1). The isolates were received as stabs, which were streaked onto LB agar plates. The following day, a swab of the growth was used to inoculate an overnight culture in LB medium, which was subsequently frozen as a master stock for each mutant strain. To verify the desired mutations, each master stock was streaked on LB agar containing 10 µg/mL tetracycline to select for cells containing the Tn5 insertion that characterizes the transposon library strains. For each, a single opaque colony was picked and restreaked onto LB agar containing tetracycline. The following day, a single opaque colony was picked and used to inoculate an overnight culture in LB medium containing 10 µg/mL tetracycline; the resulting cultures were frozen as working stocks for each mutant strain and were used for all future experiments. The desired transposon insertion was confirmed by PCR using primers that flank the proper ORF; primers are listed in Table 2. Throughout the manuscript, strains are always organized by ascending gene number, as in Figure 1. A few genes' annotated names differ from the conventional names for the given protein; these genes' annotated name and conventional name are both provided here, but the conventional names are used throughout for consistency and clarity: actP2/copA1, copA/cueO, copR/cusR, cus/cusS, actP1/copA2.

**Generation of Complemented Strains**

The previously described Tn7-based strategy of complementation was used to complement the transposon mutant strains (Kumar et al., 2010; Jacobs et al., 2014a,b). The wild-type ORF and native promoter were amplified using the primers listed in Table 2. If the ORF was not located directly downstream of the promoter due to operonic structure, the two DNA fragments were fused by SOE PCR. The full construct was amplified with Phusion Hot Start Polymerase (Thermo Fisher Scientific) and then cloned into pUC18T-mini-Tn7-hph using the added restriction enzyme sites (Table 2). The resulting plasmids were transformed into electrocompetent DH5α *E. coli* and transformants were selected for on LB agar with 100 µg/mL ampicillin. Correct plasmid construction was confirmed by digestion and then the resulting *E. coli* strains were mated with the AB5075 transposon mutant strains using the tri-parental protocol described by Jacobs et al. (2014b). Briefly, 100 µL of four overnight cultures (*E. coli* carrying the complementation construct in pUC18T-mini Tn7-hph, two helper strains, and the AB5075 mutant strain) grown in LB medium with the appropriate antibiotics were mixed, washed twice, and spotted onto plain LB agar. The mating was allowed to proceed for 1 to 24 h. The cells in the spot were collected with a sterile loop and plated on LB agar with 25 µg/mL chloramphenicol to kill the *E. coli* strains and 250 µg/mL hygromycin to select for *A. baumannii* with the Tn7 insertion. Resulting colonies were screened for Tn7 insertion using colony PCR with the previously described *att*Tn7 primers (Jacobs et al., 2014a). Single colonies were restreaked onto LB agar with 10 µg/mL tetracycline and 250 µg/mL hygromycin. A single opaque colony was used to inoculate an overnight culture in LB with tetracycline and hygromycin, which was ultimately used to create a freezer stock of the strain. The correct insertion at the *att*Tn7 site was confirmed by PCR and sequencing. All of the strains carried the Tn7 insertion in the same direction: forward, relative to *glmS*.

**Cloning of Region D From AB5075 Into Other A. baumannii Strain Backgrounds**

**Creation of pAJ100**

An *E. coli*–*A. baumannii* shuttle vector with a hygromycin resistance gene was created to move genes into multidrug-resistant *A. baumannii*. To generate this vector, pMQ300 (Kalivoda et al., 2011) was digested with BspH1 to remove the 1.7 kb region that contained yeast replication genes URA3 and CEN6/ARS4. The *A. baumannii* origin of replication of pWH1266 (Hunger et al., 1990) was amplified using the pWHori For and pWHori NcoI Rev primers described in Table 2. This insert was digested with *NcoI* and then ligated to the remaining 4.7 kb of pMQ300. The resulting 6 kb plasmid was transformed into *E. coli* and plated on LB agar with 250 µg/mL hygromycin. The resulting vector was named pAJ100. This vector has a *lacZ* insertion site for genes that contains cut sites for the following enzymes: *SmaI*, *BamHI*, *HindIII*, *KpnI*, *PstI*, and *PvuI*.

**Creation of pAJ100 – Region D**

Region D was amplified from AB5075 chromosomal DNA with Phusion Hot Start Polymerase (Thermo Fisher Scientific) using the copB complement dw and copD complement dw primers described in Table 2. The fragment was cloned into pAJ100 using the added *BamHI* restriction enzyme sites. The resulting pAJ100-Region D plasmid was electroporated into *E. coli* Top 10 cells,
### TABLE 1 | Strains used in this study.

#### Transposon mutant strains

| Strain name | Shorthand name | Locus tag | Strain name in transposon library | Tn5 location in ORF | Lab designation | Reference |
|-------------|----------------|-----------|-----------------------------------|---------------------|----------------|-----------|
| AB5075      | Wild-type      |           |                                   |                     |                |           |
| AB5075 pcoB140:T26 | pcoB:T26 | ABUW_3227 | tnaB1.kr121204p07q140            | 445 (756)           | DSM1866        | Jacobs et al., 2014a |
| AB5075 pcoA104:T26 | pcoA:T26 | ABUW_3228 | tnaB1.kr121203p02q104            | 607 (1935)          | DSM1833        | Gallagher et al., 2015 |
| AB5075 copB156:T26 | copB:T26 | ABUW_3320 | tnaB1.kr121203p06q156            | 142 (903)           | DSM1837        | Gallagher et al., 2015 |
| AB5075 copA/cueO118:T26 | cueO:T26 | ABUW_3321 | tnaB1.kr121128p02q118            | 381 (2118)          | DSM1840        | Gallagher et al., 2015 |
| AB5075 copR/cusR106:T26 | cusR:T26 | ABUW_3323 | tnaB1.kr121128p07q106            | 195 (684)           | DSM1825        | Gallagher et al., 2015 |
| AB5075 copS/cusS128:T26 | cusS:T26 | ABUW_3324 | tnaB1.kr121205p01q128            | 328 (1377)          | DSM1832        | Gallagher et al., 2015 |
| AB5075 copC160:T26 | copC:T26 | ABUW_3326 | tnaB1.kr121119p04q160            | 46 (381)            | DSM1844        | Gallagher et al., 2015 |
| AB5075 copD117:T26 | copD:T26 | ABUW_3327 | tnaB1.kr121204p06q117            | 280 (882)           | DSM1830        | Gallagher et al., 2015 |
| AB5075 ABUW_0265-193:T26 |           | ABUW_0265 | tnaB1.kr130913p10q193            | 177 (399)           | DSM1856        | Gallagher et al., 2015 |
| AB5075 czcC151:T26 |           | ABUW_0266 | tnaB1.kr121127p01q151            | 525 (1311)          | DSM1849        | Gallagher et al., 2015 |
| AB5075 czcB187:T26 |           | ABUW_0267 | tnaB1.kr121205p01q187            | 70 (1218)           | DSM1848        | Gallagher et al., 2015 |
| AB5075 czcA162:T26 |           | ABUW_0268 | tnaB1.kr121204p04q162            | 1523 (3159)         | DSM1846        | Gallagher et al., 2015 |
| AB5075 czcD174:T26 |           | ABUW_0269 | tnaB1.kr121205p08q174            | 486 (967)           | DSM1851        | Gallagher et al., 2015 |
| AB5075 ABUW_2705-139:T26 |           | ABUW_2705 | tnaB1.kr130903p04q139            | 89 (408)            | DSM1857        | Gallagher et al., 2015 |
| AB5075 cueR139:T101 |           | ABUW_2706 | tnaB1.kr130919p01q139            | 238 (402)           | DSM1858        | Gallagher et al., 2015 |
| AB5075 actP2/copA1-181:T26 |           | ABUW_2707 | tnaB1.kr121128p08q181            | 791 (2472)          | DSM1822        | Gallagher et al., 2015 |
| AB5075 ABUW_3226-176:T26 |           | ABUW_3226 | tnaB1.kr121111p04q176            | 193 (996)           | DSM1859        | Gallagher et al., 2015 |
| AB5075 ABUW_3229-182:T26 |           | ABUW_3229 | tnaB1.kr130913p04q182            | 86 (399)            | DSM1860        | Gallagher et al., 2015 |
| AB5075 ABUW_3230-184:T26 |           | ABUW_3230 | tnaB1.kr130904p01q184            | 234 (822)           | DSM1861        | Gallagher et al., 2015 |
| AB5075 ABUW_3232-131:T26 |           | ABUW_3232 | tnaB1.kr130904p04q131            | 25 (366)            | DSM1862        | Gallagher et al., 2015 |

#### Complemented mutant strains

| Strain name | Shorthand name | Lab designation | Reference |
|-------------|----------------|----------------|-----------|
| AB5075 pcoB140:T26 attTn7-hph-pcoB | pcoB:T26C | DSM1904 | This study |
| AB5075 pcoA104:T26 attTn7-hph-pcoA | pcoA:T26C | DSM1905 | This study |
| AB5075 copB156:T26 attTn7-hph-copB | copB:T26C | DSM1901 | This study |
| AB5075 copA/cueO118:T26 attTn7-hph-cueO | cueO:T26C | DSM1906 | This study |
| AB5075 copR/cusR106:T26 attTn7-hph-cusR | cusR:T26C | DSM1890 | This study |
| AB5075 copS/cusS128:T26 attTn7-hph-cusS | cusS:T26C | DSM1902 | This study |
| AB5075 copC160:T26 attTn7-hph-copC | copC:Tn5C | DSM1897 | This study |
| AB5075 copD117:T26 attTn7-hph-copD | copD:Tn5C | DSM1898 | This study |

#### Mating strains

| Strain name | Description | Lab designation | Reference |
|-------------|-------------|----------------|-----------|
| HB101/pRK2013 | Mating helper strain (oriT helper) | DSM1881 | Kumar et al., 2010 |
| DH5a/pJr/pTNS2 | Mating helper strain (transposase) | DSM1880 | Kumar et al., 2010 |
| DH5a/pUC18T mini-Tn7-hph | Tn7 with hygromycin resistance gene, hph, and MCS | DSM1883 | Jacobs et al., 2014a |
| DH5a/pUC18T mini-Tn7-hph-pcoA | Tn7 with hph and pcoA with native promoter | DSM1894 | This study |
| DH5a/pUC18T mini-Tn7-hph-pcoB | Tn7 with hph and pcoB with native promoter | DSM1895 | This study |
| DH5a/pUC18T mini-Tn7-hph-copB | Tn7 with hph and copB with native promoter | DSM1899 | This study |
| DH5a/pUC18T mini-Tn7-hph-cueO | Tn7 with hph and cueO with native promoter | DSM1893 | This study |
| DH5a/pUC18T mini-Tn7-hph-cusR | Tn7 with hph and cusR with native promoter | DSM1889 | This study |
| DH5a/pUC18T mini-Tn7-hph-cusS | Tn7 with hph and cusS with native promoter | DSM1900 | This study |
| DH5a/pUC18T mini-Tn7-hph-copC | Tn7 with hph and copC with native promoter | DSM1893 | This study |
| DH5a/pUC18T mini-Tn7-hph-copD | Tn7 with hph and copD with native promoter | DSM1896 | This study |

(Continued)
which were plated on 250 $\mu$g/mL hygromycin, 40 $\mu$g/mL X-gal, and 1 mM IPTG. A white colony was restreaked for isolation, and a single colony was used to inoculate an overnight culture of LB medium with 250 $\mu$g/mL hygromycin. Plasmid was purified and then electroporated into AB4857 and AB5711, which had been made electrocompetent using the protocol described by Jacobs et al. (2014b). Transformants were selected for on LB with 250 $\mu$g/mL hygromycin. Double purified single, opaque colonies were used to inoculate overnight cultures of LB medium with 250 $\mu$g/mL hygromycin, which were then used to create freezer stocks of the strains. Correct plasmid construction was confirmed by digestion and PCR.

**Determination of A. baumannii Sensitivity to Copper in Liquid Culture**

To assess the effect of copper on A. baumannii growth, bacterial strains were grown in the presence of increasing concentrations of copper sulfate as previously described (Williams et al., 2016; Williams and Merrell, 2019). Briefly, bacteria from overnight cultures were subcultured at an optical density (OD$_{600}$) of 0.05 in 10 mL of M9 medium containing 0.1–1.5 mM CuSO$_4$ (Aldrich, St. Louis, MO, United States) in 1 mL of CFA medium (Evans et al., 1977), and bacteria were diluted to an OD$_{600}$ of 0.05 in a large 100 mL volume. The planktonic cultures were incubated statically at 37°C for 24 h. To expose the preformed biofilms to copper, the broth was removed and was replaced with 1.5 mL of M9 medium containing 1.5 mM CuSO$_4$; biofilms were incubated for another 6 or 24 h. To determine the number of planktonic cells, at each timepoint (0, 6, and 24 h) the medium was transferred to a 1.5 mL Eppendorf tube and vortexed for approximately 5 s to disrupt any clumped bacteria in the supernatant; samples were visually inspected to confirm the absence of clumped bacteria. To determine the number of biofilm cells, the biofilm was scraped from the sides and bottom of the well with a pipette tip and was resuspended in 1 mL of PBS. Samples of both planktonic cells and biofilm cells were plated to enumerate CFU. Three biologically independent experiments were performed.

**Assessing Copper Sensitivity of A. baumannii Biofilm**

Biofilms were grown, exposed to copper, and assessed for survival as previously described with minor adjustments to the incubation time and growth media (Williams et al., 2016). Briefly, overnight cultures of A. baumannii strains were grown in CFA medium (Evans et al., 1977), and bacteria were diluted to an OD$_{600}$ of 0.05 in 24-well tissue-culture treated plates (Corning, Corning, NY, United States) in 1 mL of CFA medium. Plates were incubated statically at 37°C for 24 h. To expose the preformed biofilms to copper, the broth was removed and was replaced with 1.5 mL of M9 medium containing 1.5 mM CuSO$_4$; biofilms were incubated for another 6 or 24 h. To determine the number of planktonic cells, at each timepoint (0, 6, and 24 h) the medium was transferred to a 1.5 mL Eppendorf tube and vortexed for approximately 5 s to disrupt any clumped bacteria in the supernatant; samples were visually inspected to confirm the absence of clumped bacteria. To determine the number of biofilm cells, the biofilm was scraped from the sides and bottom of the well with a pipette tip and was resuspended in 1 mL of PBS. Samples of both planktonic cells and biofilm cells were plated to enumerate CFU. Three biologically independent experiments were performed.

**Quantification of Intracellular Copper via ICP-MS**

**Sample Collection**

To measure the intracellular copper concentration, dry cell samples were collected and analyzed by inductively coupled plasma mass spectrometry (ICP-MS). Overnight cultures were subcultured to an OD$_{600}$ of 0.05 in a large 100 mL volume. The subcultures were grown for 2.5 h to reach log phase, and a time zero (T0) sample was collected. The sample collection procedure...
### TABLE 2 | Oligonucleotides used in this study.

| Flanked gene | Annotation | Primer name | Sequence (5’–3’) |
|--------------|------------|-------------|------------------|
| ABUW_0265    |            | ABUW_0265-up | GTGCAATTTCATCAAGCACATG |
|              |            | ABUW_0265-dw | CAATATAAAGAATGCGGTGGG |
| ABUW_0266    | czcC       | czcC-up     | TGGGACATATTCTCAACC |
|              |            | czcC-dw     | GCCATCCGACCTTTAATGCT |
| ABUW_0267    | czcB       | czcB-up     | CACCATCAGCCGCGAAGAC |
|              |            | czcB-dw     | TGGCGGAAAATGCCTATTAGC |
| ABUW_0268    | czcA       | czdA-up     | GCAACATGTCGCAATTGTG |
|              |            | czdA-dw     | GCTTCTACTGAAGATGATGA |
| ABUW_0269    | czdD       | c zdD-up    | GCAACAGTGTCGCAATTGTG |
|              |            | czdD-dw     | GGCACAGTGTCGCAATTGTG |
| ABUW_0611    |            | ABUW_0611-up | GTCATTTACCGAGGCTGAC |
|              |            | ABUW_0611-dw | GCACTGTCAAAGATCGCT |
| ABUW_2705    |            | ABUW_2705-up | CATGACACAGCTTAAACCC |
|              |            | ABUW_2705-dw | CCTGTCGACAGGAGATATTAC |
| ABUW_2706    | cveR       | cveR-up     | GAGGTCGTGAGGATATCAGTCAAG |
|              |            | cveR-dw     | GTCATTGAGTTAATTGAGG |
| ABUW_2707    | copA1      | CopA2-up    | GCTTGAAGCCATCGAGT |
|              |            | CopA2-dw    | CCTGACGGATATGGCCTC |
| ABUW_3226    |            | ABUW_3226-up | GTTATTTACCGGCTGATAAAC |
|              |            | ABUW_3226-dw | GTTCTCCAGATGTGAGAGAC |
| ABUW_3227    | pcoB       | CopB2-up    | CTGAAAATGAGAAAGGAGCTAQ |
|              |            | CopB2-dw    | CATAGGCATTTTTGTGCG |
| ABUW_3228    | pcoA       | CopA2-up    | GTCGACACAAAAATTATATCC |
|              |            | CopA2-dw    | GAAAGAACGCGCAACATTC |
| ABUW_3229    |            | ABUW_3229-up | CCAAGAATAATAGTGTCTAAGG |
|              |            | ABUW_3229-dw | GAGAGGCGCGGATGGTTT |
| ABUW_3230    |            | ABUW_3230-up | GGTATACGTGTTGTAATTACATG |
|              |            | ABUW_3230-dw | CACTCTGGTGATGGTTGAAG |
| ABUW_3220    | copB       | CopB1-up    | CCTTAATCTAAGTAGAGGAG |
|              |            | CopB1-dw    | CTTTAGATAGTAGCTGTCG |
| ABUW_3221    | cveO       | ABUW_3221-up | CGGTTGCAACATAGTTAGAG |
|              |            | CopA1-dw    | GCACAAATGTCAGCTATAC |
| ABUW_3322    |            | ABUW_3322-up | CGGTTGCAACATAGTTAGAG |
|              |            | ABUW_3322-dw | GTTAAATTAGGCGTACAGAG |
| ABUW_3323    | cusR       | CusR-up     | CTTAAATAGGCTACAGGAG |
|              |            | CusR-dw     | GCATACGGGAAATGCTG |
| ABUW_3324    | cusS       | CusS-up     | GGAATGGGGCTATCCTAGAG |
|              |            | CusS-dw     | GAAAGAGCTACCTGACCTC |
| ABUW_3325    | copA2      | CopA1_up_NEW | GCCGGAGATGACCTTCAAG |
|              |            | CopA1_dw_NEW | GAGGAAGCGACAGGTATAG |
| ABUW_3326    | copC       | CopC-up     | GGAATTCTACTGGTGTATTAC |
|              |            | CopC-dw     | CTGTCGCCATAGATAGC |
| ABUW_3327    | copD       | CopD-up     | GCAATTACTAGCGCCG |
|              |            | CopD-dw     | GAAAGCATTGTCCTACTC |

**Complementation cloning primers**

| Fragment      | Primer name | Sequence (5’–3’) |
|---------------|-------------|------------------|
| pcoAB promoter| pcoAB promoter up | CCGGGTACCGggtagtgtagagagcctaa |
|              | pcoAB promoter dw-A | GGGCTGTCAATCGGTTATGAGCATTAatgtagtaggctctctactaataaa |
|              | pcoAB promoter dw-B | AGAAATAAATACATGGATGCGGTACCaatagtagtagctctctactaataaa |
| pcoA ORF      | pcoA complement up | TTTTAGTAGAAGACATGTCATTAggtctcaataaagaattagcagcc |
|              | pcoA complement dw | CCGGGAATCGagaaagcaacgcaacatatac |
| pcoB ORF      | pcoB complement up | TTTTAGTAGAAGACATGTCATTAggtctcaataaagaattagcagcc |
|              | pcoB complement dw | CCGGGAATCGagaaagcaacgcaacatatac |
| copAB promoter| copAB promoter up | CCGGGTACCGagaaagcaacgcaacatatac |
|              | copAB promoter dw | CCGGGTACCGagaaagcaacgcaacatatac |
was as follows: 6 mL of culture was pelleted in three 2-mL tubes using a 4°C table top centrifuge at max speed for 30 s; the supernatant was aspirated. The three pellets were resuspended in 1 mL of cold wash solution (PBS with 0.5 mM EDTA) in a single tube and pelleted again; the supernatant was aspirated. The wash steps were repeated for a total of three washes. After the final aspiration, the tube was left open to begin air drying. A 0.5 mL sample of the culture was separately collected in a cuvette to measure the OD

A 0.5 mL sample of the culture was collected in a cuvette to measure the OD

The following day, the samples were cooled to room temperature and diluted with 2 mL of MilliQ water, for a final HNO\textsubscript{3} concentration of 6%. Where necessary, the samples were transferred to metal-free 15-mL conical tubes for ICP-MS analysis. The metal content of the samples was then analyzed by quantitative ICP-MS. Copper concentrations were determined by injecting diluted samples into an Agilent 7700x ICP-MS (Agilent Technologies, Santa Clara, CA, United States). Copper levels were detected using an Octopole Reaction System cell (ORS) in He mode. The ICP-MS parameters used for the analysis were: an RF power of 1550 W, an argon carrier gas flow of 0.99 L/min, helium gas flow of 4.3 mL/min, octopole RF of 200 V, and OctP bias of −18 V. Samples were directly infused using the 7700x peristaltic pump with a speed of 0.1 rps and a micromist nebulizer. Copper concentrations in samples were derived from a calibration curve generated by a series of dilutions of a copper atomic absorption standard (Fluka Analytical, St. Louis, MO, United States) prepared in the same matrix as the samples. Data analysis was performed using Agilent’s Mass Hunter software (4.4 version).

**Calculations**

The value originally obtained from ICP-MS (ppb, µg/L) was divided by the molecular weight of copper (63.546 g/mol) to determine the concentration of copper in the diluted sample (µM). The sample concentration was multiplied by the total course. Three biologically independent replicates were performed for both experiments.

**Copper Measurement**

The dry cell pellets were thawed at room temperature and dissolved in 200 µL of concentrated HNO\textsubscript{3} (trace metal grade) overnight at 80°C. The following day, the samples were cooled to room temperature and diluted with 2 mL of MilliQ water, for a final HNO\textsubscript{3} concentration of 6%. Where necessary, the samples were transferred to metal-free 15-mL conical tubes for ICP-MS analysis. The metal content of the samples was then analyzed by quantitative ICP-MS. Copper concentrations were determined by injecting diluted samples into an Agilent 7700x ICP-MS (Agilent Technologies, Santa Clara, CA, United States). Copper levels were detected using an Octopole Reaction System cell (ORS) in He mode. The ICP-MS parameters used for the analysis were: an RF power of 1550 W, an argon carrier gas flow of 0.99 L/min, helium gas flow of 4.3 mL/min, octopole RF of 200 V, and OctP bias of −18 V. Samples were directly infused using the 7700x peristaltic pump with a speed of 0.1 rps and a micromist nebulizer. Copper concentrations in samples were derived from a calibration curve generated by a series of dilutions of a copper atomic absorption standard (Fluka Analytical, St. Louis, MO, United States) prepared in the same matrix as the samples. Data analysis was performed using Agilent’s Mass Hunter software (4.4 version).

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FIGURE 2 | Growth of highly copper sensitive AB5075 mutant and complementation strains. Strains were grown for 6 h at 37°C in M9 medium supplemented with the concentration of CuSO₄ indicated in the legend. Growth was measured both by turbidity (OD₆₀₀, data not shown) and enumeration of viable colonies (CFU/mL). Black lines are used to group the mutant strain with the corresponding complemented derivative. The data are presented as geometric mean and SEM of three biologically independent experiments. Two-way ANOVA with Tukey’s adjustment for multiple comparisons was used to compare growth in each copper concentration at each timepoint. The symbols indicate which copper treatments were statistically different (P < 0.05) from the 0 mM control, as follows: ~, growth in 1.5 mM CuSO₄ was different from the control; +, growth in 1 and 1.5 mM CuSO₄ was different from the control; *, growth in 0.5, 1, and 1.5 mM CuSO₄ was different from the control; and #, growth in 0.25, 0.5, 1, and 1.5 mM CuSO₄ was different from the control.

volume (2.2 mL) to determine the quantity of copper in the sample (nmol). The total amount of copper was divided by the sample volume (6 mL) to determine nmol/mL of cells and divided by the OD₆₀₀ to determine the nmol/ODU of cells. The data was plotted as nmol/ODU vs. time. This protocol was recently described in more detail as a methods chapter (Williams et al., 2019).

Infection of Galleria mellonella Caterpillars

Acinetobacter baumannii strains were grown overnight in LB medium with the appropriate antibiotics: tetracycline for transposon mutant strains, and both tetracycline and hygromycin for complemented strains. Overnight cultures were diluted to an OD₆₀₀ of 0.05 and were grown 3 h to log phase. The cells were adjusted in PBS to a specific OD₆₀₀ to achieve the desired CFU/mL, and were then diluted 1:10 in PBS with 0.01% bromophenol blue dye for visibility. For a dose of 5 × 10⁴ CFU, bacteria were adjusted to an OD₆₀₀ of 0.2–0.3, which corresponded to 10⁷ CFU/mL; for a dose of 5 × 10⁵, bacteria were adjusted to an OD₆₀₀ of 2–3, or 10⁸ CFU/mL. The exact number of bacterial cells in the inoculum was determined by serial dilution and plating for enumeration of CFU. The actual doses ranged from 3.53–9.63 × 10⁴ to 3.38–9.60 × 10⁵.

Galleria mellonella larvae (Vanderhorst Wholesale, Inc., Saint Marys, OH, United States) were utilized within 4 days of receipt. Larvae weighing 200–300 mg were used. The injections were carried out as described previously (Jacobs et al., 2014a).
FIGURE 3 | Growth of moderately copper sensitive AB5075 mutant and complementation strains. Strains were grown for 6 h at 37°C in M9 medium supplemented with the concentration of CuSO₄ indicated in the legend. Growth was measured both by turbidity (OD₆₀₀, data not shown) and enumeration of viable colonies (CFU/mL). Black lines are used to group the mutant strain with the corresponding complemented derivative. The data are presented as geometric mean and SEM of three biologically independent experiments. Two-way ANOVA with Tukey’s adjustment for multiple comparisons was used to compare growth in each copper concentration at each timepoint. The symbols indicate which copper treatments were statistically different (P < 0.05) from the 0 mM control, as follows: ~, growth in 1.5 mM CuSO₄ was different from the control; +, growth in 1 and 1.5 mM CuSO₄ was different from the control.

with minor changes. Briefly, 5 µl of the sample was injected into the last left proleg using a 10-µl glass syringe (Hamilton, Reno, NV, United States) fitted with a 31G needle. Each experiment included control groups of non-injected larvae and PBS-injected larvae. All larvae were incubated at 37°C, and death was assessed at 24 h intervals for 5 days. Larvae were considered dead if they didn’t respond to physical stimulus. All larvae that progressed to the pupation stage of the life cycle were excluded. For consistency, wild-type, mutant strain, and complemented strain infected groups that are plotted together were injected in the same experiment. Experiments were repeated using two or three different orders of larvae with 12 to 18 larvae per experimental group; the total n ranged from 25 to 48 larvae per strain. A single data set was excluded in which the wild-type strain killed only 30% of the caterpillars; no data from this set of infections was used.

Murine Pneumonia Infection Model
Mice were infected with A. baumannii as described previously (Jacobs et al., 2014a). Briefly, 6-week-old female BALB/c mice were first rendered temporarily neutropenic by intraperitoneal injection with cyclophosphamide on days −4 and −1 before infection. On day 0, mice were anesthetized with 2–5% isoflurane gas and inoculated intranasally with approximately 5 × 10⁶ CFU of A. baumannii in a total volume of 25–50 µL PBS. Animals were monitored for morbidity and mortality for 7 days, and humanely euthanized with CO₂ inhalation when necessary. All mouse studies were conducted in accordance with the Guide for the Care and Use of Laboratory
**Animals** (National Research Council, 2011), and procedures were approved by the Institutional Animal Care and Use Committee at the Walter Reed Army Institute of Research (protocol 16-BRD-488). Experiments with the *cusR* mutant strain and complemented derivative were performed in four biological replicates with five mice per group (*n* = 20 per strain), spaced two and two over time to ensure reproducibility; two biological replicates were completed with the *copD* mutant strain and complemented derivative (*n* = 10 mice per strain).

**Data Analysis and Statistics**

All graphs and statistical analyses were carried out using GraphPad Prism 7 (GraphPad Software, Inc., La Jolla, CA, United States). A two-way analysis of variance (ANOVA) with Tukey’s correction for multiple comparisons was used to evaluate differences in growth in various copper treatments. Data are presented as the geometric mean and standard error of the mean. A two-way ANOVA with Sidak’s or Dunnett’s correction was used to compare intracellular copper concentrations. Data are presented as arithmetic mean and range. An ordinary one-way ANOVA with Dunnett’s correction followed by linear contrasts with a Bonferroni adjustment was used to compare biofilm and planktonic cells percent survival at each concentration and time point to the wild-type. Data are presented as arithmetic mean and range. Survival curves were compared using the Mantel–Cox log rank test with Holm’s correction for multiple comparisons. For all tests, a two-sided α level set at 0.05 determined significance.

**RESULTS**

**Mutant Strains of AB5075 Exhibit Copper Sensitivity**

Many putative copper-related genes have been identified in *A. baumannii*; however, their predicted functions are based solely on protein homology and the role of these proteins in copper resistance is largely unknown. To investigate the contribution of individual genes to copper resistance in AB5075, we used the available arrayed transposon mutant strain library (Gallagher et al., 2015) to initiate our studies. To this end, we obtained 21 mutant strains, each of which contained a T26 transposon insertion in a putative copper resistance gene. Specifically we targeted the 22 genes found in regions A–D that we previously identified as putative copper resistance genes in AB5075 (Williams et al., 2016); only one small ORF, ABUW_2708, was not represented in the available mutant strain library. To assess the role of the remaining 21 genes in copper resistance, each of the mutant strains was individually tested in the same growth experiment. Complete restoration of copper resistance to wild-type levels was observed in the *cusR*, *cusS*, and *copD* complemented strains (Figure 2). Partial restoration of copper resistance was observed in the *pcoB*, *pcoA*, and *copC* complemented strains (Figures 2, 3). No functional complementation was observed in the *pcoA* and *pcoB* complemented strains (Figure 3); the complemented derivative of the *pcoB* mutant strain actually grew slightly less than its parent strain in 1 mM copper sulfate. The lack of complementation perhaps suggests that the identified phenotypic changes are not purely due to the transposon insertion in *pcoA* and *pcoB*, or that this complementation strategy was insufficient for these two particular genes. Taken together, our mutational analyses indicate that many of the identified putative copper resistance genes indeed contribute to copper resistance in *A. baumannii*. Moreover, the results suggest that the genes found in region D are crucial for the highest levels of copper resistance.

**Region D Enhances Copper Resistance in Other *A. baumannii* Strains**

Our previous analysis of multiple clinical isolates of *A. baumannii* identified two isolates, AB4857 and AB5711, that both showed inherently reduced copper resistance as compared to AB5075. Notably, neither of these strains carried the genes from region D in their genomes (Williams et al., 2016). Given our finding that mutations in the genes located in region D caused the greatest copper sensitivity in AB5075 (Figure 2), we hypothesized that the transfer of region D to AB5711 and AB4857 would increase their overall level of copper resistance. Indeed, when region D from AB5075 was carried on the plasmid pAJ100 in these strains, pAJ100-Region D conferred increased copper resistance comparable to the level seen in AB5075; the empty pAJ100 vector did not affect resistance (Figure 4). Thus, these results also...
indicate that the genes in region D are important for high levels of copper resistance in *A. baumannii*.

**Biofilms of Mutant Strains Retain Increased Copper Sensitivity**

Biofilm formation is an important component of the *A. baumannii* lifecycle; the bacterium is known to create biofilms on both biotic and abiotic surfaces, which aids in fomite-based transmission and in virulence (Longo et al., 2014; Zurawski et al., 2019). Moreover, biofilms are notorious for their ability to decrease the effectiveness of antibiotic treatment (Holby et al., 2010). Because any future copper-based therapeutics would likely need to be effective against *A. baumannii* found within a biofilm structure, we next tested the copper sensitivity of biofilms formed by the copper sensitive mutant strains as compared to wild-type AB5075. Biofilms were established in complex media for 24 h before switching the medium to M9 medium supplemented with 1.5 mM CuSO$_4$. CFU were enumerated from both the biofilm and supernatant at 0, 6, and 24 h post copper shock, and data were expressed as percent survival relative to T0. For the wild-type AB5075, a slight decrease in biofilm CFU was observed at 6 h, but no obvious sensitivity to copper was observed in the planktonic population at this timepoint or in the biofilm or planktonic populations at 24 h (Figure 5). While the *pcoA* and *pcoB* mutant strains of region B did not demonstrate survival differing from the wild-type strain in these biofilm conditions, the strains bearing mutations in region D (*copB, cueO, cusR, cusS, copC, copD*) each displayed statistically significant decreases in survival of both biofilm and planktonic cells relative to the wild-type strain; a 1–3 log loss in recoverable CFU was observed. The *copB* and *cueO* mutant strains displayed an early defect but recovered to wild-type levels by 24 h. Conversely, the *cusR, cusS, copC*, and *copD* mutant strains did not recover. These data demonstrate that most mutant strains retain copper sensitivity relative to the wild-type strain even within a biofilm structure, suggesting that future copper resistance targeting therapeutics would likely not be hindered by any inherent copper resistance of a biofilm.

**Copper-Sensitive Mutant Strains Accumulate More Intracellular Copper**

To begin to understand the ability of the wild-type strain to adapt to copper stress as well as the nature of the copper homeostasis defects in the copper sensitive mutant strains, we utilized ICP-MS to temporally measure copper accumulation in strains exposed to a copper shock (0.25 mM). Previous studies in other bacteria have demonstrated that intracellular copper accumulates in mutant strains bearing mutations in copper-related proteins, including *pcoB, pcoA, copA, cusR*, and *cusS* (Outten et al., 2001; Lee et al., 2002; Gonzalez-Guerrero et al., 2010; Gudipaty et al., 2012). To optimize the assay, we focused our initial efforts on wild-type AB5075 and the *cusR* mutant strain. In wild-type AB5075, the intracellular copper concentration increased ~10x one minute following copper exposure. However, the level reduced to ~5x in approximately 15 min and fully recovered to baseline levels 60 min following removal of excess copper from the media (Figure 6). Thus, in the wild-type strain, the cells appear to respond to copper stress via the deployment of efflux mechanisms that reduce the level of intracellular copper. In comparison, for the copper sensitive *cusR* mutant strain, intracellular copper levels immediately increased ~10x, but showed no reduction while copper was present. However, after copper removal, the level similarly recovered to baseline within 60 min (Figure 6). These data suggest that in the presence of excess copper, the *cusR* mutant cells are unable to control the intracellular copper concentration and cannot efflux copper ions as well as the wild-type cells.
Based on the above results, we modified the sampled timepoints and collection strategy in order to simultaneously test the wild-type and all eight of the mutant strains that previously demonstrated increased copper sensitivity (Figures 2, 3). Of note, the revised collection strategy resulted in a longer period of copper exposure as the washing steps took ~20 min longer to collect the cells. Using this procedure, the copB and cueO mutant strains each showed significantly higher accumulation of copper relative to the wild-type strain during the 30 min copper shock. In addition, the cusR, cusS, and copD mutant strain accumulated a higher concentration of intracellular copper within the additional time of copper exposure found with the modified washing steps. All strains demonstrated recovery following the removal of the copper stress (Figure 6). Taken together, these results indicate that many of the mutant strains have defects in copper homeostasis and copper efflux, which likely contribute to their copper sensitivity.

Copper Resistance Genes Are Important for Virulence in Galleria mellonella Larvae

Given that copper is an important component of immune defense, we hypothesized that copper resistance is an important trait that affects virulence of A. baumannii. To begin to test this possibility, we assessed the ability of the copper sensitive mutant strains to kill G. mellonella caterpillars. This model was chosen because: (1) G. mellonella have been established as an inexpensive and simple infection model for many pathogenic bacteria, including A. baumannii, (2) these caterpillars can be maintained at 37°C, (3) these caterpillars have both humoral and cellular immune response pathways, and (4) this invertebrate model has been used for identification of bacterial virulence factors and virulence results often correlate with those obtained in mammalian models (Peleg et al., 2009; Jacobs et al., 2014a; Tsai et al., 2016). Groups of G. mellonella were individually infected with each strain of A. baumannii, incubated at 37°C, and observed for 5 days. Consistent with previous experiments with AB5075 (Jacobs et al., 2014a), at a dose of approximately 5.0 × 10⁴ CFU, wild-type AB5075 killed ∼85% of the G. mellonella, with most death occurring in the first 2 days post-infection (Figure 7). In contrast, the copper sensitive mutant strains were all attenuated in this model and killed significantly less G. mellonella than the wild-type strain (Figure 7). The pcoB, pcoA, cueO, cusS, and copD mutant strains were the most attenuated and killed less than 30% of the G. mellonella. The copB and cusR mutant strains were moderately attenuated and killed ∼55% of G. mellonella. The copC mutant strain was very mildly attenuated and killed ∼70% of G. mellonella. To ensure that the strains from the AB5075 transposon library did not show a general defect in the G. mellonella model, we additionally selected and tested five mutant strains that did not display a copper sensitive phenotype in vitro; none of these strains were attenuated for virulence in the G. mellonella model (Supplementary Figure S5).

As with the in vitro phenotypes, we also sought to functionally complement the virulence defects seen in the G. mellonella model. However, infection with the complemented strains resulted in similar rates of G. mellonella death as those infected with the corresponding mutant strain; in some cases, the complemented strains appeared slightly more attenuated (Supplementary Figures S3, S4). Based on these data, we speculated that the insertion at the attTn7 site was causing unintended attenuation in this model; to our knowledge, no published studies have shown functional complementation using the Tn7-based strategy in assays of G. mellonella survival. To test this possibility, a strain carrying an attTn7 insertion of only the hygromycin resistance gene was created and then compared in virulence to the wild-type

![Figure 5](image_url)
May be due to disruption of an important genetic element. The mechanistic reason for this effect is unknown, but the results suggest that the inability to functionally complement the parental strains (Supplementary Figure S5) even further attenuated (Supplementary Figure S5). These conclusions are strengthened by a lack of attenuation of other transposon mutant strains that did not demonstrate increased copper sensitivity (Supplementary Figure S5), however, these conclusions would clearly be further strengthened by functional complementation in this model.

CusR and CopD Are Important for Virulence in a Murine Pneumonia Infection Model

Given the attenuated virulence observed in the G. mellonella model, we next wished to determine if A. baumannii copper resistance is also an important virulence determinant during infection of a mammalian host. To this end, we selected two of the mutant strains and their complemented derivatives to test in a murine pneumonia model; the cusR and copD mutant strains were selected due to their high level of copper sensitivity and successful in vitro complementation (Figure 2), as well as their phenotypic differences from the wild-type strain in all other assays, including virulence in G. mellonella (Figures 5–7). The murine pneumonia model was chosen because lung infections are among the most common A. baumannii infections, and AB5075 copper sensitivity has been characterized in this model (Jacobs et al., 2014a). As shown in Figure 8, the wild-type strain killed 85% of mice by 7 days post-infection. However, the cusR mutant strain was attenuated in this model; death was delayed relative to wild-type and only 65% of the mice died. Virulence of the cusR mutant strain was restored by complementation of the cusR gene in trans; 85% of the mice were killed by the complemented strain by 7 days post-infection. The copD mutant strain showed an even more dramatic virulence defect and only killed 10% of mice as compared to 90 and 60% by the wild-type and complemented strains, respectively. The ability to functionally complement both the copD and cusR mutant strain phenotypes in the pneumonia model further supports our conclusions concerning failure to achieve complementation in the G. mellonella model. In summary, both CusR and CopD individually contribute to virulence of A. baumannii in the murine pneumonia model.

DISCUSSION

Novel therapeutics are desperately needed to treat infections with drug-resistant A. baumannii and other superbugs. Because copper homeostasis is an essential process and copper ions themselves have potent toxicity, we sought to identify important copper resistance determinants in A. baumannii; we theorize that these could potentially be targeted with future therapeutics. Using a focused approach, we identified genes in A. baumannii with...
putative roles in copper resistance, and we demonstrated that 8 of 21 tested genes individually contributed to in vitro copper resistance of AB5075; mutant strains displayed significant copper sensitive phenotypes during planktonic growth (Figures 2, 3). Mutation of genes carried in region D caused the largest increase in copper sensitivity, indicating that this region is crucial for high levels of resistance. Furthermore, when a plasmid-borne copy of region D was introduced into A. baumannii strains that naturally lacked region D, copper resistance was dramatically increased (Figure 4). Moreover, strains bearing mutations of genes in region D showed enhanced copper-dependent killing even when found in a biofilm structure (Figure 5). The copper sensitivity is likely due to the fact that many of the mutant strains (copB, cueO, cusR, cusS, copD) demonstrated markedly increased accumulation of intracellular copper as compared to wild-type AB5075, which was able to efficiently efflux copper ions (Figure 6). Finally, when tested in vivo in G. mellonella and a murine pneumonia model, the copper sensitive mutant strains showed distinct attenuation in virulence. In the G. mellonella model, the mutant strains killed less G. mellonella than the wild-type strain, with the strongest attenuation observed with the pcoB, cueO, cusS, and copD mutant strains (Figure 7). Similarly, in the murine mouse model, the cusR and copD mutant strains were also attenuated and the attenuation phenotype was able to be complemented (Figure 8).

En masse, our data indicate that copper resistance is mediated by many genes in AB5075 and show that copper resistance contributes to virulence.

Though the absolute functions of the proteins encoded by each of the genes showing homology to copper related systems...
resistance. Thus, there is potential that these genes could also contribute to resistance to other metals; such possible roles warrant further study.

We utilized multiple assays to analyze in vitro copper resistance and to assess virulence. In many cases, we observed corroborating results across multiple assays. For instance, we observed similar patterns of sensitivity in our planktonic and biofilm growth assays. The four strains with the greatest increase in copper sensitivity during planktonic growth (cusR, cusS, copC, copD) also demonstrated sustained, increased copper sensitivity in the context of biofilm growth. The four additional strains that showed more moderate increases in copper sensitivity in the planktonic growth assay, displayed temporarily increased (copB, cueO) or no change in sensitivity (pcoA, pcoB) relative to the wild-type background in the context of biofilm growth. Interestingly, relative to the most copper sensitive mutant strains, the copB and cueO mutant strains seemed to be more copper sensitive in the biofilm assay than in the planktonic growth assay; the reason for this change in the pattern of sensitivity among the strains is unknown, however, different culture conditions (rich vs. minimal media) could play a role. Additionally, we did not observe a direct correlation between in vitro copper sensitivity and attenuation of virulence in the G. mellonella infection model. For example, the two most attenuated mutant strains in the G. mellonella model (pcoB and cueO) had only moderate copper sensitivity in the planktonic growth and biofilm survival assays, while the cusR and copC mutant strains demonstrated strong phenotypes in in vitro assays but were only mildly attenuated for virulence in G. mellonella. These differences among assays highlight the importance of the methods chosen to assess genes of interest; the roles of individual genes may vary depending on environmental conditions, and this will affect the experimental outcomes and the conclusions drawn. Indeed, experimental differences may account for contrasting results in this work and those published recently by another group (Alquethamy et al., 2019). Under the conditions we tested, we did not observe increased copper sensitivity of either of the copper ATPase mutant strains (Supplementary Figure S1) or attenuation in the G. mellonella infection model (Supplementary Figure S5).

We originally hypothesized that perhaps CopA1 and CopA2 were functionally redundant, resulting in an inability to observe a copper-sensitive phenotype. However, a recently published study of the copper ATPases in AB5075 found that the gene referred to as copA1 in our publication (ABUW_2707, region B), but not copA2 (ABUW_3325, region D), contributed to copper resistance (Alquethamy et al., 2019); those results suggest that the CopA1 and CopA2 proteins are not functionally redundant. While the exact reason for the difference in study results is not clear, it is worth noting that the experimental methods used were different, e.g., growth medium (which dramatically affects free copper concentrations) and infection model; additionally, the various utilized mutant strains differed in their site of transposon insertion and transposon orientation, which may cause differential polar effects. Given the disparity in the results of the studies, it is currently unclear what the individual roles of these copper ATPases are in A. baumannii, and additional studies will be required to clearly define the role of these factors in copper resistance.
Our studies identified many copper resistance proteins that individually contributed to virulence in the chosen models; these proteins have a variety of putative functions including copper efflux, oxidation, chaperoning, and regulation. Copper-related genes with similar functions have been shown to be important for virulence in many bacterial and fungal pathogens including *Pseudomonas aeruginosa*, *Pseudomonas fluorescens*, *Xanthomonas campestris* pv. *campestris*, *Xanthomonas citri* subsp. *citri*, *Streptococcus pneumoniae*, *Streptococcus mutans*, *Salmonella Typhimurium*, *Mycobacterium tuberculosis*, *Neisseria gonorrhoeae*, *Listeria monocytogenes*, *E. coli*, *Staphylococcus aureus*, *Campylobacter jejuni*, *Aspergillus fumigatus*, and *A. baumannii* (Francis and Thomas, 1997; Schwan et al., 2005; Zhang and Rainey, 2007; Hu et al., 2009; White et al., 2009; Achard et al., 2010; Gonzalez-Guerrero et al., 2010; Osman et al., 2010; Ward et al., 2010; Corbett et al., 2011; Hsiao et al., 2011; Shafeeq et al., 2011; Wolschendorf et al., 2011; Yan and Wang, 2011; Djoko et al., 2012; Marrero et al., 2012; Chaturvedi et al., 2014; Shi et al., 2014; Subashchandrabose et al., 2014; Johnson et al., 2015; Garcia et al., 2016; Park et al., 2017; Wiemann et al., 2017; Gardner and Olson, 2018; Purves et al., 2018; Zapotoczna et al., 2018; Alquethamy et al., 2019). Of note, three types of copper-related proteins that have been previously shown to contribute to virulence of other bacteria were found to also contribute to virulence of *A. baumannii*: copper ATPases, copper oxidases, and copper chaperones. In the majority of the previous studies, the mutated copper resistance gene of interest was a copper ATPase, homologous to *copA1* and *copA2* of AB5075. Of note, a role for *CopA1* in virulence of *A. baumannii* was recently described (Alquethamy et al., 2019). However, we did not observe copper sensitivity or reduced virulence in either of our copper ATPase mutant strains (Supplementary Figures S1, S5). Copper oxidases have been shown to be important for virulence of *S. pneumoniae*, *C. jejuni*, and *S. aureus* (Achard et al., 2010; Gardner and Olson, 2018; Zapotoczna et al., 2018). Here, we demonstrated that both copper oxidases of AB5075, PcoA and CueO, contribute to virulence of *A. baumannii*. Similarly, chaperone proteins
have been previously linked to virulence in *S. pneumoniae* and *S. mutans* (Johnson et al., 2015; Garcia et al., 2016), and we found that the putative chaperone CopC contributes in *A. baumannii*. Though their functions may be similar, it is unclear how related CopC of *A. baumannii* is to the chaperones of other bacteria, as these chaperone proteins are small and have minimal sequence identity or similarity. While these three types of proteins are interesting as putative drug targets, an important consideration in identifying novel drug targets is assessment of homology with host proteins and potential toxicity from off-target activity. Copper ATPase and oxidase proteins have amino acid homology to human proteins, and the copper ATPases are also functionally similar. Thus, designing therapeutics that target these functions may be complicated by homology to the essential human proteins, e.g., human copper ATPases ATP7A and ATP7B. While not insurmountable, homology to host proteins certainly makes design of novel therapeutics more challenging.

Of particular interest, we identified a number of copper-associated proteins that have not previously been shown to contribute to virulence in other bacterial species. To our knowledge, this is the first publication to report attenuated virulence in any bacterial species of strains lacking the putative transporters PcoB, CopB, or CopD (Wolschendorf et al., 2011; Subashchandrabose et al., 2014). We also observed attenuated virulence of the strains bearing mutations in either portion of the two-component system CusRS. To our knowledge, this is the first report that this copper-sensing two-component system contributes to virulence of a bacterial species during mammalian infection. Additionally, none of these proteins share amino acid sequence homology with human proteins, and therefore may be targeted more readily.

While the focus of this work was copper resistance in *A. baumannii*, we note that copper homeostasis mechanisms are well-conserved across bacterial species; the open reading frames studied here were originally identified in AB5075 due to their conserved predicted protein sequence. Therefore, we expect that any novel therapeutics that are targeted against these proteins may have broad-spectrum antibacterial activity. Despite the fact that individual proteins are well-conserved, the presence of these proteins or their homologs varies across bacterial species and even amongst strains (Hernandez-Montes et al., 2012). Indeed, we observed that not all the same copper resistance proteins are present in all clinical isolates of *A. baumannii*. Thus, these differences will be an important consideration when choosing which copper resistance genes may be the best therapeutic targets. Given all of these possible targets, copper homeostasis components could provide an attractive starting place for the development of future antimicrobial therapies.

The immune system has harnessed the antimicrobial power of copper for use against pathogens and copper-mediated immune mechanisms are known to play a role in clearance of bacterial pathogens (Sheldon and Skaar, 2019). Because the innate immune response of phagocytic cells is crucial for clearance of *A. baumannii* infection, we hypothesize that copper resistance contributes to virulence of this pathogen by enhancing immune evasion and survival *in vivo*. Therefore, we predict that novel therapeutics targeting copper resistance of bacterial pathogens would enhance immune clearance by reducing bacterial survival in the copper-rich phagosomal compartment. The role of copper in the immune system has been demonstrated by prior studies that utilized copper-replete and copper-deficient cell lines and animal models. For example, macrophages that are pre-incubated with copper are able to kill more internalized bacteria than controls (White et al., 2009). Conversely, copper-deficient macrophages or macrophages that are unable to mobilize copper to the phagosomal compartment due to loss of function of ATP7A are less efficient at killing bacterial pathogens (White et al., 2009; Achard et al., 2012; Ladomersky et al., 2017). The importance of copper in immunity has also been demonstrated on an organismal level. Animals with copper-deficient diets are more susceptible to infection, while those with copper-rich diets more efficiently clear infection (Newberne et al., 1968; Wolschendorf et al., 2011; Gardner and Olson, 2018). Furthermore, mice with a myeloid-specific knockout of ATP7A, the copper ATPase that pumps toxic copper ions into the macrophage phagosome, are more susceptible to infection with *Salmonella Typhimurium* (Ladomersky et al., 2017). Comparable future studies using *A. baumannii* will help to shed light on whether copper resistance of this organism aids in virulence by enabling evasion of host copper-mediated immune strategies.

In summary, we identified several additional copper resistance proteins that may serve as potential therapeutic targets, including copper transporters PcoB, CopB, and CopD, chaperone CopC, oxidases CueO and PcoA, and regulatory proteins CusR and CusS. Future work using targeted antibodies and small molecules will seek to determine the efficacy of targeting these copper resistance proteins as a successful treatment strategy. Given the relative conservation of these proteins and pathways among bacteria, we believe several are viable therapeutic targets to treat *A. baumannii* infection as well as other multidrug-resistant bacterial pathogens, for which novel treatments are urgently needed.

**DATA AVAILABILITY STATEMENT**

The datasets generated for this study are available on request from the corresponding author.

**ETHICS STATEMENT**

The animal study was reviewed and approved by the Institutional Animal Care and Use Committee at the Walter Reed Army Institute of Research.

**AUTHOR CONTRIBUTIONS**

CW, SM, DZ, and DM conceived and designed the experiments. CW, HN, YA, RR, AJ, SS, and RA-T performed the experiments. CW and DM wrote the manuscript.
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SUPPLEMENTARY MATERIAL
The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fmicb.2020.00016/full#supplementary-material

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Conflict of Interest: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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