Population Structure, Antibiotic Resistance, and Uropathogenicity of *Klebsiella variicola*

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ABSTRACT *Klebsiella variicola* is a member of the *Klebsiella* genus and often misidentified as *Klebsiella pneumoniae* or *Klebsiella quasipneumoniae*. The importance of *K. pneumoniae* human infections has been known; however, a dearth of relative knowledge exists for *K. variicola*. Despite its growing clinical importance, comprehensive analyses of *K. variicola* population structure and mechanistic investigations of virulence factors and antibiotic resistance genes have not yet been performed. To address this, we utilized *in silico*, *in vitro*, and *in vivo* methods to study a cohort of *K. variicola* isolates and genomes. We found that the *K. variicola* population structure has two distant lineages composed of two and 143 genomes, respectively. Ten of 145 *K. variicola* genomes harbored carbapenem resistance genes, and 6/145 contained complete virulence operons. While the β-lactam *bla*<sub>LEN</sub> and quinolone *oqxAB* antibiotic resistance genes were generally conserved within our institutional cohort, unexpectedly 11 isolates were nonresistant to the β-lactam ampicillin and only one isolate was nonsusceptible to the quinolone ciprofloxacin. *K. variicola* isolates have variation in ability to cause urinary tract infections in a newly developed murine model, but importantly a strain had statistically significant higher bladder CFU than the model uropathogenic *K. pneumoniae* strain TOP52. Type 1 pilus and genomic identification of altered *fim* operon structure were associated with differences in bladder CFU for the tested strains. Nine newly reported types of pilus genes were discovered in the *K. variicola* pan-genome, including the first identified P-pilus in *Klebsiella* spp.

IMPORTANCE Infections caused by antibiotic-resistant bacterial pathogens are a growing public health threat. Understanding of pathogen relatedness and biology is imperative for tracking outbreaks and developing therapeutics. Here, we detail the phylogenetic structure of 145 *K. variicola* genomes from different continents. Our results have important clinical ramifications as high-risk antibiotic resistance genes are present in *K. variicola* genomes from a variety of geographic locations and as we demonstrate that *K. variicola* clinical isolates can establish higher bladder titers than *K. pneumoniae*. Differential presence of these pilus genes in *K. variicola* isolates may indicate adaption for specific environmental niches. Therefore, due to the potential of multidrug resistance and pathogenic efficacy, identification of *K. variicola* and *K. pneumoniae* to a species level should be performed to optimally improve patient outcomes during infection. This work provides a foundation for our improved understanding of *K. variicola* biology and pathogenesis.

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Klebsiella variicola was initially believed to be a plant-associated, distant lineage of *Klebsiella pneumoniae*; however, it has subsequently been recovered from human clinical specimens (1). Despite increasing knowledge on the distinctness of *K. variicola*, *K. pneumoniae*, and *Klebsiella quasipneumoniae*, misidentification within the clinical microbiology lab commonly occurs (2, 3). This may have clinical implications, as one study demonstrated that *K. variicola*-infected patients have higher mortality than *K. pneumoniae*-infected patients (4). Furthermore, several virulence genes (VGs), including siderophores, allantoin utilization genes, and glycerate pathway genes, have been reported in select *K. variicola* strains (5, 6). *K. variicola* has been shown to contain a large pan-genome that is distinct from *K. quasipneumoniae* and *K. pneumoniae*, but the functional consequences of differential gene content have not been explored (2, 7).

In this study, we retrospectively analyzed a cohort of *Klebsiella* isolates collected from 2016 to 2017 at Washington University in St. Louis School of Medicine/Barnes-Jewish Hospital Clinical Microbiology Laboratory (WUSM) for possible *K. variicola* strains using matrix-assisted laser desorption ionization–time of flight mass spectrometry (MALDI-TOF MS) and yggE PCR-restriction fragment length polymorphism (RFLP) assays. We performed Illumina whole-genome sequencing (WGS) to compare *K. variicola* from our institution with publicly available genomes in the first global evaluation of this species. We particularly focused on annotation of canonical *Klebsiella* VGs and antibiotic resistance genes (ARGs) and then assessed their functional consequences using *in vitro* assays and *in vivo* murine infections. Our results demonstrate that population structure, antibiotic resistance, and uropathogenicity of *K. variicola* are generally similar to *K. pneumoniae*, but variability among *K. variicola* genomes has important clinical implications with various strain efficacies in a murine model of urinary tract infection (UTI).

### RESULTS

Average nucleotide identity and MALDI-TOF MS can differentiate *K. variicola* from *K. pneumoniae*. We performed Illumina WGS on 113 isolates that are commonly misidentified as *K. pneumoniae* (*K. variicola* [n = 56], *K. quasipneumoniae* [n = 3], *K. pneumoniae* [n = 53], and *Citrobacter freundii* [n = 1]). They were identified by Bruker Biotyper MALDI-TOF MS and yggE RFLP assays from a variety of adult infection sites (see Table S1 in the supplemental material). The isolates were retrieved from the Barnes-Jewish Hospital clinical microbiology laboratory (St. Louis, MO, USA) in 2016 to 2017. We used pyANI with the mummer method to calculate the pairwise average nucleotide identity (ANI) between the isolates in our cohort and retrieved publicly available *Klebsiella* genomes (n = 90) (8, 9) (Table S1). The *C. freundii* isolate was originally classified as *K. pneumoniae* from the Vitek MS MALDI-TOF MS v2.3.3 but was later determined to be *Citrobacter freundii* by Bruker Biotyper MALDI-TOF MS. The yggE PCR-RFLP was indeterminate for this isolate. Confirmatory yggE PCR-RFLP had 94.6% (53/56) concordance with MALDI-TOF for prediction of *K. variicola* within our cohort (Fig. 1). While one genome was dropped from downstream analysis, the other 55 WUSM *K. variicola* genomes all had >95% ANI with the reference genome of *K. variicola* At-22 (5). *K. variicola* HKUOPOLA (GCA_001278905.1) had >95% ANI with *K. quasipneumoniae* ATCC 7000603 reference genome but not *K. variicola* At-22, indicating that it is likely a misannotated *K. quasipneumoniae* isolate and not a *K. variicola* isolate. The remainder of the NCBI *K. variicola* genomes clustered with *K. variicola* At-22 and the WUSM *K. variicola* cohort. One hundred percent (41/41) of the *K. pneumoniae* genomes from NCBI that were suspected to be *K. variicola* due to BLAST similarity had >95% ANI with *K. variicola* At-22 but not *K. pneumoniae* HS11286 or *K. pneumoniae* CAV1042 (Fig. 1).

Hierarchical clustering of the pairwise ANI values replicated previous phylogenetic analysis showing that *K. pneumoniae* and *K. quasipneumoniae* are more closely related to each other than to *K. variicola* (Fig. 1). Interestingly, the clustering pattern within *K.
*K. variicola* indicated that two isolates, KvMX2 (FLLH01.1) and YH43 (GCF_001548315.1), are more closely related to one another than to the remainder (143/145) of the *K. variicola* genomes. Given that *K. quasipneumoniae* can be differentiated into two subspecies based on ANI with the BLAST method (ANIb), we used the JSpecies ANIb program to specifically compare KvMX2 and YH43 with *K. pneumoniae* ATCC BAA-1705, *K. quasipneumoniae* ATCC 700603, and 3 other *K. variicola* genomes (10). KvMX2 and YH43 have 98.02% ANIb with one another but an average of 96.67%, 96.65%, and 96.68% ANIb with WUSM_KV_53, WUSM_KV_15, and *K. variicola* At-22, respectively (Table S1). Consistent with our pyANI ANIm result, none of the *K. variicola* strains had ≥95% ANIb with *K. pneumoniae* ATCC BAA-1705 or *K. quasipneumoniae* ATCC 700603. These data suggest that MALDI-TOF MS or *yggE* PCR-RFLP may be effective means to differentiate *K. variicola* from *K. pneumoniae* in the absence of WGS.

*K. variicola* population structure has 2 lineages and 26 clusters in the second lineage. Core-genome alignment of the 1,262 genes at 90% identity shared by strains in all *Klebsiella* species and a *Kluyvera georgiana* outgroup shows that the *K. variicola* isolates are in a cluster with *K. pneumoniae*, *K. quasipneumoniae*, and the newly described *K. quasivariicola* (11) (Table S2; Fig. S1). Core-genome alignment of the 3,430 core genes at 95% nucleotide identity for the entire gene length by all 145 *K. variicola* genomes indicates that KvMX2 and YH43 are distantly related to the other 143 genomes (Fig. 2a; Table S2). These other genomes form a star-like phylogeny showing deep-branching clusters radiating from the center of the tree. FastGear, which uses hierBAPS to identify lineages and then searches for recombination between lineages, supported the differentiation of KvMX2 and YH43 into a separate lineage from the other genomes and identified 6 instances of recombination between these two lineages (Table S3) (12, 13).

Phylogenomic network analysis and quantification of recombination from parSNP showed minimal recombination within the 143 *K. variicola* lineage 2 genomes, with approximately 1.62% of the *K. variicola* genome believed to be recombinant (Fig. S2a; Table S3) (14). The Nearest Neighbor network of the 3,496 genes shared by the lineage 2 genomes and a recombination-free phylogenetic tree of the 143 genomes from...
FIG 2  Population structure of *K. variicola* genomes. (a) Approximate-maximum-likelihood tree of the total 145 *K. variicola* genomes and annotation of FastGear lineage identification. (b) Recombination-free parSNP tree of the closely related lineage 2 genomes with quantitative clustering from ClusterPicker (Continued on next page)
parSNP showed many deep-branching clades with a star-like phylogeny (Fig. 2b; Table S2). This tree topology was similar with and without recombination, which suggests that *K. variicola* lineages emerged early from a single common ancestor into equally distant clades across different environments (Fig. S2b). Quantitative clustering of the 143 genomes in the second lineage with ClusterPicker showed that 56.6% (81/143) of genomes fall into 26 clusters, with 57.7% (15/26) of the clusters containing more than 2 genomes (Fig. 2b) (15). Only 46.2% (12/26) of clusters contain isolates from both WUSM_KV and NCBI. The largest clusters, 24 and 21, each contain 7 genomes. Cluster 21 contained WUSM_KV_10 and 6 genomes from an analysis of patient isolates at an intensive care unit in Seattle, WA (USA). Although they were in the same cluster, WUSM_KV_10 differed from these isolates at 1,882 sites across the 4,867 genes shared at 95% identity (Table S2 and Table S4).

To better understand the context of the 4 groups in lineage 2, we aligned the 2,932 genes shared among the 145 *K. variicola* genomes, *Klebsiella* (formerly *Enterobacter*) aerogenes KCTC 2190, *K. quasipneumoniae* ATCC 700603, and *K. pneumoniae* ATCC BAA-1705 at ≥90% identity to create a dendrogram (Fig. 2c; Table S2). This method preserved the conservation of the lineage 2 groups but showed a different order. The only discrepancy observed is that, in the lineage 2 phylogenetic tree, cluster 3 appeared to be in the A group; however, both 521_SSON and 524_SBOY are more similar to C group genomes in the dendrogram. This incongruence is consistent with cluster 3 radiating away from cluster 4 near the center of the phylogenetic tree (Fig. 2b).

Addition of metadata onto the dendrogram showed that the *K. variicola* cohort spans most geographic locations, with the notable exception of Africa and Oceania (Fig. 2c). The *K. variicola* genomes showed a remarkable level of source diversity, with representative isolates from animals (n = 4), fungi (n = 2), plants (n = 7), water (n = 3), and industrial waste (n = 6). However, as a testament to the pathogenic potential of *K. variicola*, 79.5% (114/145) of genomes came from sites associated with humans. Of the human-associated sites, 40.4% (46/114) came from urine and 19.2% (22/114) came from respiratory specimens. In addition to the near-total conservation of lineage 2 across different environments (Fig. S2b), quantitative clustering suggests that *K. variicola* has a diverse population structure and can be found in a variety of environmental and host niches.

**Acquired ARGs and VGs are not restricted to any *K. variicola* cluster.** We applied ResFinder to determine the burden of acquired ARGs among the *K. variicola* strains (Fig. 3a; Table S5) (16). β-Lactamase genes were the most abundant ARG in the *K. variicola* cohort (n = 26). As expected, *blaLEN* was almost universally conserved, as 837_KPNE was the only isolate without one identified. Ten different *blaLEN* alleles were found. *blaLEN-16* was most common (51/145), followed by *blaLEN-24* (40/145) and *blaLEN-2* (31/145). Carbapenemases were rare, but *blaKPC-2* (4/145), *blaKPC-6* (1/145), *blaNDM-1* (1/145), *blaNDM-9* (3/145), and *blaOXA-48* (1/145) were each identified across a total of 10/145 strains. *blaCTX, blaSHV, blaTEM* and noncarbapenemase *blaOXA* genes were also identified, but we did not detect any class C β-lactamase genes or non-*blaNDM* class B β-lactamase genes. Aminoglycoside ARGs (n = 10), including members of the *aac, aad, aph,* and *str* families, comprised the second most abundant class. ARGs against folate synthesis inhibitors (n = 8), quinolones (n = 7), amphenicols (n = 4), tetracyclines (n = 2), macrolides/linosamides/streptogramins (n = 2), and fosfomycin (n = 1) were also found (Fig. 3a). In addition to the near-total conservation of *blaLEN*, the quinolone efflux

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**FIG 2 Legend (Continued)**

added as alternating teal and brown labels adjacent to cluster number (1 to 26). Bootstrap support values below 80% are depicted as node labels. (c) Monophyletic groups of these clusters were colored if they were similar in the dendrogram showing the evolutionary context of the cluster compared to *K. pneumoniae* (KP), *K. quasipneumoniae* (KQ), and *K. aerogenes* (KA). Relevant metadata for initial identification, geographic location, source of isolation, and body site are adjacent to the assembly names. Bootstrap support values below 80% are depicted as node labels.
pump components *oqxAB* were found in almost all isolates (139/145). Across the 145 genomes, the median and mode number of ARGs were both 3. A 6.89% (10/145) proportion of genomes harbored ≥10 ARGs, including WUSM_KV_55 from our cohort.

We used the *K. pneumoniae* BIGSdb database ([https://bigsdb.pasteur.fr/klebsiella/klebsiella.html](https://bigsdb.pasteur.fr/klebsiella/klebsiella.html)) and BLASTN to identify canonical *Klebsiella* VGs in the *K. variicola* strains (Fig. 3b; Table S5). In contrast to ARGs, previously characterized *Klebsiella* VGs were found only sporadically in the *K. variicola* cohort. Interestingly, the *all* allantoin utilization operon and *arc, fdrA, gci, glxKR, hyi,* and *ybbWY* genes were found in the distantly related YH43 genome as well as the closely related BIDMC90, k385, and WUSM_KV_03

**FIG 3** Distribution of acquired antibiotic resistance and virulence genes in the *K. variicola* cohort. Presence/absence matrix of ARGs (a), virulence genes (b), and plasmid replicons (c) ordered for all *K. variicola* genomes against the dendrogram from Fig. 2c.
found to harbor additional quinolone (the lowest number of acquired ARGs, as it lacked ARGs. Interestingly, it possesses these isolates. We transurethrally inoculated C3H/HeN mice with 10^7 CFU/ml of 5

Given that 70% (39/56) of our cohort strains are susceptible to most antibiotics. Co-constructed a network diagram of ARGs and isolates to identify connectivity within the K. variicola cohort strains are isolated from 5 hospitals in the St. Louis region (Missouri, USA), based on first isolate per patient per year, revealed that, in general, K. pneumoniae (n = 1,522) had decreased susceptibility to all reported antimicrobials compared to K. variicola (n = 144), except for meropenem (99% susceptibility for both species). Most notably, K. pneumoniae exhibited decreased susceptibility, compared to K. variicola, with ampicillin-sulbactam (63% versus 93% susceptible), nitrofurantoin (66% versus 86% susceptible), and trimethoprim-sulfamethoxazole (80% versus 90% susceptible).

Changes in fim operon are associated with uropathogenicity in a murine UTI model. Given that 70% (39/56) of K. variicola strains from our cohort were isolated from the human urinary tract, we wanted to assess uropathogenicity in a diverse subset of these isolates. We transurethrally inoculated C3H/HeN mice with 10^7 CFU/ml of 5 individual K. variicola strains, or the model uropathogenic K. pneumoniae TOP52 strain, for comparison (Fig. 5a) (3, 18, 19). Similarly to previously published infections with K. pneumoniae TOP52, the K. variicola strains exhibited large variations in bacterial CFU recovered from the bladder at 24 h postinfection (hpi). Compared to TOP52, WUSM_KV_39 was the only isolate with a significantly increased bladder burden (P = 0.0094). Bacterial loads of WUSM_KV_10 and WUSM_KV_39 were both significantly higher than WUSM_KV_09 and WUSM_KV_14 (Fig. 5a). Despite this variability among
FIG 4  WUSM K. variicola strains have a low burden of ARGs and are generally susceptible to antibiotics. (a) Network diagram depicting each WUSM_KV isolate and ARG as nodes. ARGs are colored in accordance with predicted phenotypic resistance from ResFinder, and WUSM_KV genomes are colored by the burden of ARGs. (b) Scatter plots depicting Kirby-Bauer disk diffusion size (mm) from phenotypic susceptibility testing. Each plot represents an isolate, and the plots are colored according to CLSI interpretation. Those with atypical resistance are listed by name with putative ARGs.
FIG 5  Changes in fim operon are associated with outcomes in mouse UTI model. (a) CFU/bladder and CFU/kidney of K. pneumoniae TOP52 and WUSM KV isolates 24 h after transurethral bladder inoculation of C3H/HeN mice. Short bars represent geometric means of each group, and dotted lines represent limits of detection. (b) fimS phase assay and quantification with respective bands indicating the “ON” and “OFF” position labeled. (c) Immunoblot for FimA and GroEL, with quantification shown below. (d) Easyfig illustration of genes in the fim operon and Jalview of the nucleotides and amino acids for the fimB/fimE intergenic region and fimD gene.
bladder CFU results, the results of kidney titer determinations at 24 hpi were not significantly different among strains by ANOVA ($P = 0.1270$). As observed in the bladder, however, WUSM_KV_10 and WUSM_KV_39 achieved significantly higher kidney CFU than WUSM_KV_14.

Given the variation in bladder burden, we wanted to assess if differences in uropathogenicity could be related to expression of type 1 pili, a key virulence factor for UTI encoded by the fim operon (19, 20). In K. pneumoniae and Escherichia coli, expression of type 1 pili is controlled by a region of invertible DNA (fimS site) (20, 21). Orientation of the fimS site in the “ON” position enables production of type 1 pili and increased urovirulence. Under identical growth conditions, WUSM_KV_39 had a higher population with the fimS promoter region in the “ON” orientation than the other strains tested (Fig. 5b). Furthermore, consistent with its success in the bladder, WUSM_KV_39 was found to produce the greatest amount of FimA (the main structural component of type 1 pili), as measured by immunoblotting (Fig. 5c). WUSM_KV_03, WUSM_KV_09, and WUSM_KV_39 all produced significantly more FimA than K. pneumoniae TOP52. Interestingly, WUSM_KV_14 did not produce appreciable levels of FimA by this assay (Fig. 5c).

As we discovered significant variability in type 1 piliation, we specifically investigated changes in fim operon sequence between these isolates by viewing the Prokka coding sequence annotation in Easyfig and Jalview (Fig. 5d) (22, 23). We found that WUSM_KV_14 had a predicted truncated FimD usher sequence. A guanine-to-adenine single nucleotide polymorphism (SNP) in the fimD gene changed a predicted tryptophan residue into a premature stop codon, likely explaining the observed lack of production of type 1 pili. Additionally, in WUSM_KV_39, Prokka annotated a hypothetical protein in the intergenic region between fimB and fimE and included a gap replacing a thymine and a guanine-to-adenine SNP. The altered fimB/fimE intergenic region in WUSM_KV_39 may play a role in its increased expression of type 1 pili. Together, these data demonstrate that variation exists among K. variicola genomes that may account for differential urinary tract niche proclivity among isolates.

**K. variicola contains both conserved and novel usher genes.** The fim operon is one of the best-characterized chaperone-usher pathways (CUPs); given the observed importance of the fim operon in K. variicola uropathogenicity, we searched the pan-genome of our K. variicola cohort to identify the complete repertoire of CUP operons (24). Seventeen unique usher sequences at 95% identity were identified across the 55 WUSM K. variicola genomes, and an amino acid sequence alignment showed that they were distributed in 5 Nuccio and Baumler (25) clades (Fig. 6a; Table S6). From this analysis, we discovered 9 new usher genes previously undescribed in Klebsiella, which we name kva through kvi (Table S6). KviA and KveB usher sequences were found to cluster within the pi ($\pi$) clade, making them the first description of a P-pilus apparatus in Klebsiella. The recently named $\gamma^*$ subclade contained the greatest amount (7/17) of K. variicola usher sequences; 5 of these 7 were previously reported in K. pneumoniae, while KvcC and KvdB are first reported here.

FimD and the usher sequences for KpaC, KvaB, KpeC, and KpjC were present in all 55 WUSM K. variicola isolates (Fig. 6b). KvgC, KvhC, KviA, and KpcC were each found in only one isolate. KpgC, MrkC, KvbC, KpbC, KvcC, KveB, KvfC, and KvdB can be considered accessory usher sequences in this cohort, as they were absent in certain strains. The most notable pattern evident from the hierarchical clustering of the presence/absence for all usher genes in our K. variicola cohort is that isolates WUSM_KV_10 through WUSM_KV_21 all carry the KvdB sequence but not KpbC.

Eight of the 9 newly described usher sequences had highest BLASTP hits of $\approx 99\%$ identity across the entire length of the gene against the nonredundant protein sequences database in April 2018, and all of them were previously annotated as being found in Enterobacteriaceae, Klebsiella, or K. variicola (Table S6). All of the usher genes except kvi were in operons that included a chaperone, at least one subunit, and a putative adhesin (Table S6). KvhC, the usher protein with the lowest BLASTP identity
value, had 76% identity to several genes from Enterobacter species (Fig. S3a). The contig with the kvh operon also contained several genes that had possible roles in prophage integration and transposase activity (Fig. S3b). Our results indicate that *K. variicola* strains harbor a diverse set of usher genes, which may augment *K. variicola* fitness across a variety of environmental niches, and these operons may be acquired from other *Enterobacteriaceae*.

**DISCUSSION**

A previous phylogenomic study used split-network analysis to demonstrate that the *K. variicola* phylogroup (formerly KPIII) is distinct from *K. pneumoniae* (KPI) and *K. quasipneumoniae* (KPII) (26). As an orthogonal method, we used ANI software, the gold standard for *in silico* species delineation, to recreate this differentiation of phylogroups as separate species (8). Historically, differentiation between *K. pneumoniae* and *K. variicola* has been difficult, as evidenced by misannotation of *K. variicola* as *K. pneu-
moniae in public genome sequence databases (Fig. 1). These misannotated K. variicola strains came from a variety of geographic regions and were not exclusive to any cluster. Within our sequenced cohort, differentiation of K. variicola from K. pneumoniae and K. quasipneumoniae using MALDI-TOF MS and yggE PCR-RFLP was supported by ANI. This indicates that yggE PCR-RFLP (3) would be a feasible alternative for clinical labs across the globe lacking access to MALDI-TOF MS or WGS. Additionally, hierarchical clustering of the ANI values and core-genome phylogeny demonstrated that 2 K. variicola genomes were distinctly separate from the other 143 in our cohort. ANIb values between these genomes and the other K. variicola genomes were ~96%, similar to what was observed for K. quasipneumoniae. The differences in ANIb values contributed to the delineation of K. quasipneumoniae into two subspecies, Klebsiella quasipneumoniae subsp. quasipneumoniae and Klebsiella quasipneumoniae subsp. similipneumoniae (27). Further phenotypic comparisons, including the sole carbon source utilization used for differentiation of the K. quasipneumoniae subspecies, between KvMX2/Yh43 and other K. variicola isolates is required to unequivocally qualify these as separate subspecies (27).

Numerous studies have shown that K. pneumoniae has a deep-branching phylogenetic structure with minimal recombination occurring within K. pneumoniae strains and between K. pneumoniae and K. variicola/K. quasipneumoniae (26, 28). Importantly, though, large-scale recombination events may be clinically relevant, as evidenced by research on the origin of the frequently carbapenem-resistant ST258 lineage (29, 30). Our results demonstrate that like K. pneumoniae, K. variicola shows minimal recombination within its genome, and its population structure is composed of numerous clades in a star-like phylogeny. A star-like population structure with deep-branching relationships between isolates (n = 29 and n = 28) was also found in two previously published K. variicola phylogenetic trees (2, 31).

Similarly to our work, a previous investigation did not identify any geographic distinction when genomes from within the United States were compared to those from outside the United States (2). The 6 genomes in cluster 21 with WUSM_KV_10 were from ICU patient samples in Seattle, WA, which provides the first evidence of clonal groups responsible for K. variicola infections in some settings (32). Although they were closely related compared against all K. variicola genomes, there were still 1,882 SNPs between WUSM_KV_10 and the other 6 genomes. Interestingly, clusters were not restricted to human infections, as cluster 24 contains 3 genomes from bovine mastitis (NL49, NL58, and NL58) and hospital isolates (VRCO00246, VRCO00242, VRCO00244, and VRCO00243) (https://www.ncbi.nlm.nih.gov/bioproject/361595) (33).

As expected for K. variicola, bla LEN β-lactamases were the most conserved ARGs. A previous report unexpectedly found a K. variicola isolate that harbored the bla KCP gene commonly found in K. quasipneumoniae; however, we did not identify such instances within our cohort (2). Although chromosomally carried in K. pneumoniae, fosA was identified in only 1/145 of the K. variicola genomes (34, 35). Additionally, as previously found in K. pneumoniae clinical isolate cohorts, we found oqxAB efflux pump genes widespread across K. variicola genomes (36–38). Although these genes may be ubiquitous in K. variicola, 0 of 55 isolates we tested had resistance to ciprofloxacin; the single example with intermediate susceptibility carried a qnrB6 gene. This is not atypical for Enterobacteriaceae possessing oqxAB, as one study found 100% prevalence of oqxAB in K. pneumoniae but no quinolone resistance (37). It is possible that for K. variicola, similarly to K. pneumoniae, high expression of oqxAB is essential for phenotypic resistance to quinolones (36). In K. pneumoniae, expansion of clonal groups is associated with carbapenemase carriage (i.e., ST258 and bla KPC); however, we did not observe any associations between carbapenemase genes and K. variicola clusters. Indeed, only 1.81% (1/55) of K. variicola strains within our institutional cohort had a carbapenemase gene and the regional resistance rate for meropenem between K. pneumoniae and K. variicola in 2017 was similar. bla NDm-positive K. variicola strains have been identified in clinical and environmental samples, but bla KPC-positive genomes came exclusively from clinical sources. KPN1481 (bla NDm-1) was annotated as a urine-derived isolate, but GJ1,
K. pneumoniae is the second leading cause of urinary tract infections (45). Given previous misclassification of K. variicola as K. pneumoniae and the high frequency at which K. variicola was isolated from the urinary tract, we were interested in comparing the uropathogenicity of our K. variicola isolates to the well-studied model K. pneumoniae TOP52 isolate (3, 18, 19). We identified strain-dependent virulence capacity, with UTIs from WUSM_KV_39 yielding statistically significant higher bladder CFU than K. pneumoniae TOP52. Quantification of metrics used to study uropathogenicity in E. coli and K. pneumoniae show increased fimS in the “ON” orientation and increased FimA production by WUSM_KV_39; these findings provide a plausible explanation for why WUSM_KV_39 performed better than K. pneumoniae TOP52 and all WUSM_KV isolates excluding WUSM_KV_10 (46). While we do not yet understand the role of the putative protein identified between recombinases fimB and fimE in WUSM_KV_39, one could postulate that this difference may affect fimbral expression. Additionally, the poorest performer in the urinary tract, WUSM_KV_14, encodes a mutation resulting in a truncated fimD usher sequence which likely explains its lack of FimA production. As with other bacterial pathogens, it is likely that specific virulence factors are required for K. variicola competency in distinct body niches (47, 48). Further work is therefore warranted to test if yersiniabactin and allantoin utilization promote lung and liver infections, respectively, in K. variicola as they do in K. pneumoniae (49–52).

K. variicola carries usher genes previously identified in K. pneumoniae and 9 novel ushers (53). Interestingly, KveB and KviA are the first report of π usher proteins in Klebsiella. The best-studied π operon, pap in E. coli, is a major contributor to pyelonephritis as the PapG adhesin can bind Gal-(1-4)-Gal exposed on human kidney cells (54). Other usher genes have been shown to be essential for biofilm formation, plant cell adhesion, and murine gut colonization, further demonstrating their role in niche differentiation (53). Clustering of the presence/absence of these ushers showed the absence of KpbC but presence of KvdB in 11 of the WUSM_KV genomes, a phenomenon similar to that observed for UshC and YraJ in E. coli (55). All 4 of these usher types were found in the γγ′ clade, suggesting an exclusionary form of functional redundancy between usher genes (55). Usher genes and CUP operons are frequently exchanged horizontally between Enterobacteriaceae genera (55). Indeed, we have found that the KvhC usher protein has only 76% amino acid identity to any existing proteins in the nonredundant protein sequence database and that the kvh operon is situated next to multiple prophage- and transposase-associated genes.

In this investigation, we used phenotypic and genomic analyses to better understand the diversity of K. variicola genomes, both from our institution and across the globe (using publicly available NCBI genomes). Then, we assessed the functional consequences of ARGs and VGs toward antibiotic resistance and uropathogenicity. One limitation of our study is that our mouse infections and phenotypic analyses are performed with nonisogenic strains. If existing genetic modification systems in K. pneumoniae are shown to be useful for gene knockouts in K. variicola, further work can be performed to mechanistically validate our findings. An additional limitation is that ~30 genomes of K. variicola have been uploaded to NCBI since we initiated our comparison. These may further elucidate differences in population structure, although even with almost 300 genomes, one study indicates that K. pneumoniae diversity remains undersampled (26).

Our work represents the first large-scale genomic analysis of K. variicola across multiple institutions and the first use of a murine model to study K. variicola pathogenesis. We unequivocally show that whole-genome comparisons can separate K.
varicola from K. pneumoniae and offer convenient alternative methods for laboratories without access to WGS to differentiate these species. Importantly, we demonstrate that high-risk ARGs and VGs are present in K. varicola genomes from a variety of geographic locations. This may have clinical ramifications, as we demonstrate that some K. varicola clinical isolates can be superior uropathogens compared to K. pneumoniae. Similarly to E. coli and K. pneumoniae, the diversity of CUP operons in these isolates could complement additional acquired virulence genes and enable infection of specific niches. Therefore, it is imperative that K. varicola and K. pneumoniae continue to be differentiated in the clinical laboratory, so that we may apply data on differential gene repertoire, clinical behavior, and niche specificity to the goal of ultimately improving patient outcomes.

MATERIALS AND METHODS

Clinical Klebsiella collection. One hundred thirteen clinical Klebsiella species isolates recovered in the Barnes-Jewish Hospital microbiology laboratory (St. Louis, MO) from 2016 to 2017 were evaluated in this study. Of these, 56 were consecutively collected isolates identified by Bruker Biotyper MALDI-TOF MS as K. varicola (research-use-only database v6). This identification was confirmed using a PCR-restriction fragment length polymorphism (RFLP) assay targeting the yggE gene (F: 5’-TGTACTAAATGCCTTA CGGG-3’; R: 5’-CAGGATCTGCAAAAGTCTACT-3’; restriction enzyme: BciVI) that was designed to distinguish K. varicola from K. pneumoniae. A 94.6% proportion (53/56) of isolates were confirmed as K. varicola using the yggE PCR-RFLP assay.

The remaining 58 isolates were randomly selected from a banked collection of K. pneumoniae strains historically recovered from clinical specimens (29 from urine, 25 from blood, and 1 each from abdominal wound, tracheal aspirate, bronchial washing, and bile). Each of these isolates underwent Bruker MALDI-TOF MS and yggE PCR-RFLP to confirm their identification. Five percent (3/58) were confirmed as K. varicola using MALDI-TOF MS and the yggE PCR-RFLP assay.

Illumina whole-genome sequencing and publicly available Klebsiella genomes. Pure frozen stocks of the presumptive 113 Klebsiella isolates were plated on blood agar to isolate single colonies. Approximately 10 colonies were suspended using a sterile cotton swab into water, and total genomic DNA was extracted using the Bacteremia kit (Qiagen). An 0.5-ng amount of DNA was used as input for sequencing libraries using the Nextera kit (Illumina) (56). Libraries were pooled and sequenced on an Illumina NextSeq 2500 high-output system to obtain ~2.5 million 2 × 150-bp reads. Demultiplexed reads had Illumina adapters removed with Trimmomatic v.36 and decontaminated with DeconSeq v0.4.3 (57, 58). Draft genomes were assembled with SPAdes v3.11.0, and the scaffolds.fastas were used as input for QUAST v 4.5 to measure the efficacy of assembly (see Table S1 in the supplemental material) (59, 60). All contigs of ≥500 bp in length were annotated for open reading frames with Prokka v1.12 (61).

To increase the number of genomes for downstream analysis, 50 K. varicola genomes were obtained from NCBI genomes (https://www.ncbi.nlm.nih.gov/genome/) in September 2017 (Table S1). Additionally, as it is possible that previously sequenced K. varicola may be incorrectly described as K. pneumoniae, we submitted the complete genome of the K. varicola reference strain AT-22 to NCBI BLASTN against the nonredundant nucleotide collection and the whole-genome shotgun sequence databases using default settings in September 2017. Using this method, we obtained 41 genomes of K. pneumoniae with the minimum observed query length of 38% at 99% identity (Table S1). Given that the cohort of genomes analyzed in our study includes isolates initially misannotated, we refer to them as either the NCBI genome or assembly (https://www.ncbi.nlm.nih.gov/assembly) accession key. Sequenced and acquired isolates were analyzed using a variety of computational programs (Text S1). In silico sequence typing was performed using mlst v2.11 (https://github.com/tseemann/mlst) and the BIGSdb database (https://bigsdb.pasteur.fr/klebsiella/klebsiella.html).

Antimicrobial susceptibility testing. K. varicola isolates underwent antimicrobial susceptibility testing per laboratory standard operating procedures using Kirby-Bauer disk diffusion on Mueller-Hinton agar (BD BBL, Mueller-Hinton II agar), in accordance with Clinical and Laboratory Standards Institute (CLSI) standards. Disk diffusion results were interpreted using CLSI Enterobacteriaceae disk diffusion breakpoints (62). Briefly, 4 to 5 colonies from pure isolates were used to create a 0.5 McFarland suspension of the organism in sterile saline. A sterile, nonsterile cotton swab was dipped into the bacterial suspension, and a lawn of the organism was plated to Mueller-Hinton agar. Antimicrobial Kirby-Bauer disks were applied, and the plate was incubated at 35°C in room air for 16 to 24 h. The diameters of the zones of growth inhibition surrounding each antimicrobial disk were recorded in millimeters.

Mouse urinary tract infections. Bacterial strains from our K. varicola cohort and K. pneumoniae TOP52 were used to inoculate 7- to 8-week-old female C3H/HeN mice (Envigo) by transurethral catheterization as previously described (18, 19, 63). The K. varicola strains were selected to encompass a range of genetically distinct isolates. WUSM_KV_03 and WUSM_KV_10 were specifically chosen as they contain the all and ybt operons, respectively. Static 20-ml cultures were started from freezer stocks, grown in Luria-Bertani (LB) broth at 37°C for 16 h, and centrifuged for 5 min at 8,000 × g, and the resultant pellet was resuspended in phosphate-buffered saline (PBS) and diluted to approximately 4 × 10^7 CFU/ml. Fifty milliliters of this suspension was used to infect each mouse with an inoculum of 2 × 10^7 CFU/ml. Inocula were verified by serial dilution and plating. At 24 hpi, bladders and kidneys were aseptically harvested, homogenized in sterile PBS via Bullet Blender (Next Advance) for 5 min, serially
diluted, and plated on LB agar. All animal procedures were approved by the Institutional Animal Care and Use Committee at Washington University School of Medicine.

**Phase assays.** To determine the orientation of the *fimS* phase switch in *Klebsiella*, a phase assay was adapted as previously described (20). An 817-bp fragment including *fimS* was PCR amplified using *Taq* polymerase (Invitrogen) and the primers 5′-GGGACAGATACGCGTTTGAT-3′ and 5′-GGCCTAACCTGAACGGTTTGA-3′ and then digested with HinfI (New England Biolabs). Digestion products were separated by electrophoresis on a 1% agarose gel. A phase-ON switch yields products of 605 and 212 bp, and a phase-OFF switch yields products of 496 and 321 bp.

**FimA and GroEL immunoblots.** Acid-treated, whole-cell immunoblotting was performed as previously described using 1:2,000 rabbit anti-type 1 pilus and 1:500,000 rabbit anti-GroEL (Sigma-Aldrich) primary antibodies (64, 65). Amersham ECL horseradish peroxidase-linked donkey anti-rabbit IgG (GE Healthcare) secondary antibody (1:2,000) was applied, followed by application of Clarity enhanced chemiluminescence (ECL) substrate (Bio-Rad Laboratories). The membrane was developed and imaged using a ChemiDoc MP Imaging System (Bio-Rad Laboratories). Relative band intensities were quantified using Fiji (https://fiji.sc/) (66).

**Statistics.** CFU/bladder and CFU/kidney for both experimental replicates were used as input for ordinary one-way ANOVA to judge significance. Pairwise comparisons of CFU/bladder and CFU/kidney values were performed by using the nonparametric Mann-Whitney U test. Similarly, normalized quantifications of relative FimA amounts (FimA/GroEL) and *fimS* in “ON” position (*fimS* “ON”/*fimS* “OFF”) were compared using the Mann-Whitney U test. All *P* values of <0.05 were considered significant, and all calculations were performed in GraphPad Prism v7.04.

**Accession number(s).** The genomes have all been deposited in NCBI under BioProject accession no. PRJNA473122.

### SUPPLEMENTAL MATERIAL

Supplemental material for this article may be found at [https://doi.org/10.1128/mBio.02481-18](https://doi.org/10.1128/mBio.02481-18).

- **TEXT S1**, DOCX file, 0.04 MB.
- **FIG S1**, TIF file, 1.5 MB.
- **FIG S2**, TIF file, 3.4 MB.
- **FIG S3**, TIF file, 1 MB.
- **TABLE S1**, XLSX file, 0.1 MB.
- **TABLE S2**, XLSX file, 0.6 MB.
- **TABLE S3**, XLSX file, 0.02 MB.
- **TABLE S4**, TXT file, 0.1 MB.
- **TABLE S5**, XLSX file, 0.04 MB.
- **TABLE S6**, XLSX file, 0.1 MB.

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