A Klebsiella variicola Plasmid Confers Hypermucoviscosity-Like Phenotype and Alters Capsule Production and Virulence

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Hypermucoviscosity (hmv) is a capsule-associated phenotype usually linked with hypervirulent Klebsiella pneumoniae strains. The key components of this phenotype are the RmpADC proteins contained in non-transmissible plasmids identified and studied in K. pneumoniae. Klebsiella variicola is closely related to K. pneumoniae and recently has been identified as an emergent human pathogen. K. variicola normally contains plasmids, some of them carrying antibiotic resistance and virulence genes. Previously, we described a K. variicola clinical isolate showing an hmv-like phenotype that harbors a 343-kb pKV8917 plasmid. Here, we investigated whether pKV8917 plasmid carried by K. variicola is linked with the hmv-like phenotype and its contribution to virulence. We found that curing the 343-kb pKV8917 plasmid caused the loss of hmv, a reduction in capsular polysaccharide ($P < 0.001$) and virulence. In addition, pKV8917 was successfully transferred to Escherichia coli and K. variicola strains via conjugation. Notably, when pKV8917 was transferred to K. variicola, the transconjugants displayed an hmv-like phenotype, and capsule production and virulence increased; these phenotypes were not observed in the E. coli transconjugants. These data suggest that the pKV8917 plasmid carries novel hmv and capsule determinants. Whole-plasmid sequencing and analysis revealed that pKV8917 does not contain mmpA2 genes; thus, an alternative mechanism was searched. The 343-kb plasmid contains an IncFIB backbone and shares a region of ~150 kb with a 99% identity and 49% coverage with a virulence plasmid from hypervirulent K. variicola and multidrug-resistant...
INTRODUCTION

Hypermucoviscosity (hmv) is a distinctive trait displayed by hypervirulent *Klebsiella pneumoniae* (hvKpn), and it is visualized by the formation of a viscous filament >5 mm by strains grown on agar plates. However, quantitative tests are recommended to distinguish hmv from non-hmv strains (Walker and Miller, 2020). A recent study showed that capsule and hmv are not the same process; hence, the capsule serves as a scaffold for hmv (Walker et al., 2019). The molecular mechanism underlying hmv in hvKpn strains involves multiple global regulators acting directly or indirectly on the cps region (Ares et al., 2015; Walker and Miller, 2020). Among the set of proposed regulators, the *rmpADC* locus is highly correlated with hmv in hvKpn (Walker et al., 2019; Walker et al., 2020). However, it is not well known whether hmv could contribute to a higher virulence level when hypervirulence traits are absent. Different works have reported *K. pneumoniae*, *K. variicola* and *K. quasipneumoniae* isolates displaying the hmv-like phenotype as the genes involved are unknown and apparently not related to *rmpADC* (Yu et al., 2006; Li et al., 2014; Arena et al., 2015; Cubero et al., 2016; Garza-Ramos et al., 2016; Harada et al., 2019; Imai et al., 2019). Consequently, this scenario suggests the existence of *rmpADC/rmpA2*-independent hmv-like pathways in *Klebsiella*.

*Klebsiella variicola* has gained clinical recognition as an emerging human pathogen not only for its capacity to acquire antimicrobial resistance and virulence genes but also for being widespread and types of infections (Rodriguez-Medina et al., 2019). We previously described the first *K. variicola* clinical isolate categorized as hypermucoviscous that harbors one 343-kb plasmid (Garza-Ramos et al., 2015; Rodriguez-Medina et al., 2019). The present study focuses on the genetic analysis of the pKV8917 plasmid as a determinant capable of transfer hmv-like and its role in virulence.

MATERIALS AND METHODS

Bacterial Isolate Identification

*Klebsiella variicola* 8917 clinical isolate was previously reported (Garza-Ramos et al., 2015). Briefly, the isolate was obtained from the sputum of an elderly male patient at the Hospital Regional Centenario de la Revolución Mexicana in Morelos, Mexico, in 2011.

*K. pneumoniae*. The pKV8917-unique region harbors a cellulose biosynthesis cluster (bcs), fructose- and sucrose-specific (*fru/scr*) phosphotransferase systems, and the transcriptional regulators *araC* and *iclR*, respectively, involved in membrane permeability. The hmv-like phenotype has been identified more frequently, and recent evidence supports the existence of *rmpADC/rmpA2*-independent hmv-like pathways in this bacterial genus.

Conjugation Assays, *in vitro* Induction of Nalidixic Acid-Resistant *Klebsiella variicola* Mutants and Plasmid Stability

The non-hmv *K. variicola* F2R9 (DSM 15968, ATCC BAA 830) nalidixic acid-resistant and *Escherichia coli* J53-2 rifampicin-resistant were used as recipient strains. The conjugation experiments were carried out in broth. *In vitro* nalidixic acid resistance was induced in *K. variicola* F2R9 because this strain is susceptible to all antibiotic families (except ampicillin). A nalidixic acid-resistant mutant was obtained from a serial passage of the susceptible strain in a medium with increasing concentrations of nalidixic acid (from 16 to 128 µg ml⁻¹). Briefly, an overnight culture of *K. variicola* F2R9 lacking nalidixic acid was diluted 1:100 and then inoculated in fresh medium containing 16 µg ml⁻¹ nalidixic acid. Thereafter, bacteria from the previous stage were serially passaged daily in Luria-Bertani broth (LB) containing increasing concentrations of nalidixic acid. Finally, the bacterium that grew in 128 µg ml⁻¹ was considered a mutant, and this mutant was used as a recipient for conjugation experiments. The donor and recipient strains were cultured to logarithmic phase [optical density (OD) of approximately 0.6] at 37°C with shaking. Next, 200 µl of donor cells and 800 µl of recipient cells were mixed, placed into a new tube, and then incubated at 33°C overnight. After incubation, the mixture was serially diluted. Potential transconjugants were selected on LB agar plates containing 100 µg ml⁻¹ rifampicin and 16 µg ml⁻¹ potassium tellurite K₂TeO₃ for *E. coli* J53-2 transconjugants, and for *K. variicola* F2R9 transconjugants, plates contained 128 µg ml⁻¹ nalidixic acid and 16 µg ml⁻¹ potassium tellurite K₂TeO₃. The transconjugants were confirmed to have indistinguishable enterobacterial repetitive intergenic consensus (ERIC) patterns to the parent strains (Versalovic et al., 1991). Additionally, for *K. variicola* transconjugant analysis, we selected three genes of the *K. variicola* multilocus sequence typing scheme (Barrios-Camacho et al., 2019) that had a different allele compared with to *K. variicola* 8917. The *leuS*, *pgi*, and *pyrG* genes were selected for PCR amplification and Sanger sequencing (see Supplementary Table 2). Later, a phylogenetic analysis of the concatenated sequence of these three genes was performed. In addition, to confirm the acquisition of the plasmid, we amplified the plasmid markers *terW*, *fruA*, and *scrK* by PCR, and plasmid profile determination was performed according to the alkaline lysis method described by Kieser (1984). The *E. coli* NCTC 50192 strain, which contains 154-, 66-, 48-, and 7-kb plasmids, was used as a molecular size
1 ml of absolute ethanol to 250 µl of bacterial cultures was mixed with 100 µl of 1% zwittergent in 100-mM citric acid. Then, the mixtures were dissolved in 200-µl phosphate-buffered saline (PBS) to an OD600 of 250. The suspensions were centrifuged for 5 min at 8,000 × g, and the bacterial pellets were resuspended in 1 ml of PBS containing potassium tellurite (16 µg ml−1). Colonies that were unable to grow in LB supplemented with potassium tellurite were considered as possible plasmid-free cells (cured strain). Plasmid-free cells were confirmed by analyzing the plasmid profile, amplification of the terW, fruA, and scrK genes, and ERIC-PCR to confirm the genetic relationship of the cured strain and the parent strain. Plasmid-free cells were grown in MacConkey agar to evaluate the hypermucoviscous phenotype by string test and mucoviscosity assay.

Plasmid Curing

Plasmid curing was performed according to El-Mansi et al. (2000). A solution of sodium dodecyl sulfate (SDS; 10% w/v, pH 7.4) was added to 2 × LB medium to give final concentrations of 0.5, 1, 2, 3, 4, and 5% SDS. An inoculum of 100 µl of a preculture of the tested strain was added to the SDS-containing LB and incubated at 27°C with shaking. Separated colonies were obtained by plating 100 µl of the cells in SDS-containing LB. Later, separated colonies were replicated on LB agar and LB agar containing potassium tellurite (16 µg ml−1). Colonies that were unable to grow in LB supplemented with potassium tellurite were considered as possible plasmid-free cells (cured strain). Plasmid-free cells were confirmed by analyzing the plasmid profile, amplification of the terW, fruA, and scrK genes, and ERIC-PCR to confirm the genetic relationship of the cured strain and the parent strain. Plasmid-free cells were grown in MacConkey agar to evaluate the hypermucoviscous phenotype by string test and mucoviscosity assay.

Mucoviscosity Assay

In addition to the qualitative string test, the sedimentation degree was assessed according to Bachman et al. (2015). An overnight culture was pelleted by centrifugation at 8,000 × g and resuspended in phosphate-buffered saline (PBS) to an OD600 of ∼1. The suspensions were centrifuged for 5 min at 1,000 × g, and the OD600 of the supernatants was measured. Final readings were normalized to the OD600 of the wild-type culture before centrifugation. The results are presented as the mean and standard deviation of the data of three experiments. K. variicola F2R9 and E. coli J53-2 were used as a negative control for comparing K. variicola 8917, K. variicola F2R9, and E. coli J53-2 transconjugants, and as a positive control, the strain K. pneumoniae 10271 rmpA+/KL2 (Catalan-Najera et al., 2019) was used.

Extraction and Quantification of Capsule

Glucuronic acid (GA) content was extracted and quantified using a colorimetric assay, as previously described (Lin et al., 2009). Briefly, 500 µl of bacterial cultures was mixed with 100 µl of 1% zwittergent in 100-mM citric acid. Then, the mixtures were incubated at 50°C for 20 min and centrifuged 5 min at 8,000 × g; the capsular polysaccharide (CPS) was precipitated by adding 1 ml of absolute ethanol to 250 µl of supernatants. Pellets were dissolved in 200-µl water, and then, 1,200 µl of 12.5-mM borax concentrated in H2SO4 was added. Mixtures were vigorously vortexed, boiled for 5 min, and cooled. Twenty microliters of 0.15% 3-hydroxydiphenol in 0.5% NaOH solution was added to the mixture, and the absorbance was measured at 520 nm. The GA content was determined from a standard curve of GA and expressed in microgram/10⁹ CFU.

Phagocytosis Resistance Assay

The phagocytosis assay was performed as described elsewhere (Ares et al., 2016). In brief, THP-1 (ATCC TIB-202) human monocytes (differentiated to macrophages with 200-nM phorbol 12-myristate 13-acetate for 24 h; 6 × 10⁵) were seeded into 24-well tissue culture plates. Bacteria were grown in 5 ml of LB to the exponential phase. Macrophages were infected at a multiplicity of infection of 100 in a final volume of 1-ml Roswell Park Memorial Institute 1640 tissue culture medium supplemented with 10% heat-inactivated fetal bovine serum. To synchronize the infection, plates were centrifuged at 2,000 × g for 5 min. Plates were incubated at 37°C under a humidified 5% CO2 atmosphere. After 2 h, cells were rinsed three times with PBS and incubated for an additional 60 min with 1 ml of Roswell Park Memorial Institute 1640 containing 10% fetal bovine serum and gentamicin (100 µg ml−1) to eliminate extracellular bacteria. Cells were then rinsed again three times with PBS and lysed with 0.1% Triton X-100. After homogenization, 10-fold serial dilutions were plated onto LB agar plates to determine total CFUs.

Serum Killing Assay

Human blood was obtained from healthy individuals, and resistance to serum killing was performed as previously described (Hughes et al., 1982; Podschun et al., 1991). An inoculum of 25 µl of bacterial suspension (∼10⁶ CFU) prepared from the mid-log phase was mixed with 75 µl of pooled human serum. Viable counts were checked at 0, 1, 2, and 3 h of incubation at 37°C. Each strain was tested three times, and the mean results were expressed as a percent of surviving inoculum. The response to serum killing in terms of viable counts was scored using six grades classified as serum sensitive (grade 1 or 2), intermediate sensitive (grade 3 or 4), or serum resistant (grade 5 or 6). Inactivated serum at 56°C was used as control.

Mouse Infection Model

We implemented healthy and diabetic models. Healthy male BALB/c mice were obtained from the animal facility of the National Institute of Public Health at the age of 6–7 weeks old. Diabetic mice were induced through two doses of intraperitoneal injection of the β-pancreatic cell toxin streptozotocin; the first dose was 75 mg/kg, and the second dose was 135 mg/kg. Last, glucose was measured after 6 days of post-induction, and we considered diabetic conditions > 200 mgdl⁻¹ glucose. K. variicola 8917, F2R9, and ΔpKV8917 were inoculated into healthy and diabetic mice as follows. Bacterial cell cultures were centrifuged at 10,000 × g for 5 min. The supernatants were discarded and the bacterial pellets resuspended in 1 ml of PBS solution. The solution containing PBS and bacterial cells was diluted, and 100 µl containing 1 × 10⁵ to 8 × 10⁵ was injected via the intraperitoneal route in healthy and diabetic mice. Animals were monitored twice daily for 10 days of post-inoculation. The transconjugant F2R9_TC14 was inoculated as described earlier in healthy mice. Animals were monitored twice daily for 10 days of post-inoculation, after which mice were killed.
studies were approved by the Biosafety Committee at the National Institute of Public Health.

Whole-Genome Sequencing of Klebsiella variicola 8917

The sequence of the complete nucleotide sequence from the chromosome and plasmid of K. variicola 8917 was determined using long (MiniION) and short-read (Illumina MiSeq) sequencing. For MiSeq sequencing, genomic DNA was extracted using the DNeasy Blood and Tissue Kit (QIAGEN Inc., Germany). For MiniION sequencing, genomic DNA was extracted using the blood and cell culture kit. DNA concentrations were measured using the Qubit dsDNA HS Assay Kit (Fisher Scientific Inc.) on a Qubit 3.0 fluorometer (Fisher Scientific Inc.). De novo hybrid assembly of the short Illumina reads and long MiniION reads was performed using Canu v2.0 and SPAdes v3.1.1 assemblers (Bankevich et al., 2012; Koren et al., 2017). Chromosome and plasmid were annotated under National Center for Biotechnology Information’s Prokaryotic Genome Annotation Pipeline. The sequence type of the K. variicola genomes included in this study was determined according to the K. variicola multilocus sequence typing platform. The identification of virulence genes and plasmid replicon type was determined by the BLASTn search and PlasmidFinder 2.1 tools found in the Center for Genomic Epidemiology.

Comparative Genome and Plasmid Analysis

Plasmid and genome comparisons with other structures were generated by BLAST Ring Image Generator v.0.95.22 (Alikhan et al., 2011) using 90 and 70% as the upper and minimum thresholds, respectively. We conducted a genome comparison between three K. variicola genomes: 8917, TUM111415 (accession no. BI0K01000005.1), and KvL18 (accession no. PRJNA612181). The plasmid comparison was carried out by aligning the plasmids pKV8917, p15WZ-82_Vir (accession no. CP032356.1), pVir_030666 (accession no. CP027063.3), and pKSBl_10f unnamed2 (accession no. CP024517.1) and the chromosomes of Raoultella terrigena strain NCTC13098 (accession no. NZ_LR131271.1), Pantoea caeleiphila (accession no. NZ_PDET01000010.1), and Kosakonia radicincitans strain DSM 16656 (accession no. CP018016.1). The complete sequences of the chromosome and plasmid of K. variicola 8917 were updated in the GenBank database (CP063403 and CP063404).

Identification of Capsule Mutations

Capsular types were determined from genome sequence data based on a complete K locus primary reference database of the Kaptive v.0.5.1 (Wyres et al., 2016). The nucleotide sequence of the capsule operon of K. variicola 8917 was used as a query in BLASTn to look for strains with the same capsule type. K. pneumoniae strain QMP (accession no. LT174583.1), K. variicola 13450 (accession no. CP026013.1), and E. coli strain CSF3273 (accession no. CP026932.2) were identified as capsule-type KL114. Each protein sequence in the cps (comprising GalF and Ugd) was used to align with CLUSTALW to look for mutations in the capsule biosynthesis genes of the K. variicola 8917 genome. A phylogenetic tree was constructed with the concatenated amino acid sequence of the capsule cluster using MEGAX software. Easyfig (Sullivan et al., 2011) was used to compare the capsule genes.

Statistical Analysis

For significant differences, we implemented an unpaired two-sided Student’s t-test and one-way analysis of variance. For survival curves, two-tailed Mann–Whitney U-tests were performed using Prism 8.0 software (GraphPad Software Inc., San Diego, CA, United States).

RESULTS

Molecular and Genetic Characteristics of the Klebsiella variicola 8917 Isolate

Klebsiella variicola 8917 clinical isolate was previously identified by the string test, and a viscous filament greater than 5 cm long was observed; thus, it was considered hypermucoviscous (Garza-Ramos et al., 2015). Subsequently, a ~200-kb plasmid designated pKV8917 was determined by the Kaiser method (Rodriguez-Medina et al., 2019). Nonetheless, plasmid assembly showed a larger plasmid of 343 kb that did not carry any known antimicrobial resistance genes and encoded tellurium resistance (Rodriguez-Medina et al., 2019). Core virulence factors were predicted from the genome sequence, including the iron transport system Kfu (kfuABC), the siderophore enterobactin (entB), and the type 3 fimbriae (mrkABCDFHII). This strain encoded neither colibactin, aerobactin, nor yersiniabactin. However, we found the receptor of aerobactin (iutA; Martinez-Romero et al., 2018; Rodriguez-Medina et al., 2019). Further analysis of the implications of carrying siderophore receptors for iron scavenging is required.

pKV8917 Plasmid Confers Hypermucoviscosity-Like in Klebsiella variicola but Not in Escherichia coli

Mating experiments using K. variicola F2R9 and E. coli J53-2 as recipient strains showed that the hmv-like could be transferred to a non-hmv K. variicola but not to E. coli (Table 1). The plasmid markers terW, scrK, and fruA were amplified in the transconjugant, and resistance to tellurite was observed, confirming that the transconjugants acquired the plasmid (Table 1). In addition, the ERIC profile and the concatenated sequence of the leuS, pgi, and pyrG genes of F2R9 transconjugants were identical to the F2R9 parent strain (Supplementary Figures 1, 2).

1http://mlstkv.insp.mx
2https://cge.cbs.dtu.dk/services/PlasmidFinder/
Likewise, the ERIC profiles of the E. coli transconjugants were identical to those of the recipient strain J53-2 (Supplementary Figure 1).

The conjugation frequency was low (10^{-8} CFU) in K. variicola F2R9; in contrast, when using E. coli J53-2, the conjugation frequency was 10^{-2} CFU (Table 1). Plasmid stability was tested in the transconjugants by the amplification of plasmid markers and plasmid extraction using alkaline lysis. pKV8917 was maintained after 10 serial passages even without potassium tellurite (data not shown).

Interestingly, the K. variicola F2R9 transconjugants were string test positive; meanwhile, the E. coli J53-2 transconjugants showed a negative string test result (Table 1). However, the string displayed by the F2R9_TC5 and F2R9_TC14 transconjugants was not the same size as that displayed by the K. variicola 8917 strain. Thus, hmv-like is conferred by the pKV8917 plasmid. Considering the high frequency observed with E. coli J53-2, a second mating was assayed using J53-2_TC7 and J53-2_TC5 as donors of pKV8917 with K. variicola F2R9 as the recipient; however, this mating was unsuccessful.

### pKV8917 Plasmid Curing Resulted in the Loss of Hypermucoviscosity-Like

Bacterial cells recovered with 5% SDS were selected for further analysis. One colony (ΔpKV8917 strain) was successfully cured from a total of 1,000 colonies analyzed. This was evidenced by (i) the loss of tellurite resistance, (ii) the absence of plasmid markers terW, scrK, and fruA (Table 1), and (iii) the absence of plasmid pKV8917 determined by the Kieser protocol (Supplementary Figure 3). Interestingly, growth on MacConkey agar showed the inability of the ΔpKV8917 strain to display the viscous filament, and mucoviscosity assay reflects a slight reduction in OD measurement (Figure 1A). The loss of hmv-like was observed at a frequency of 0.1%. This result confirms the relationship that exists between hmv and the pKV8917 plasmid.

### Klebsiella variicola 8917 Is Hypermucoviscous but Sediments at Low-Speed Centrifugation

The mucoviscosity assay has been recommended as a quantitative indicator of hmv. The tested strain is subjected to low centrifugation, and the supernatant is measured at OD 600. Usually, hmv strains have poor sedimentation (Walker and Miller, 2020). Although K. variicola 8917 is hypermucoviscous, it sedimented well at low-speed centrifugation (Figure 1A). When comparing the cured (ΔpKV8917) and transconjugant (F2R9_TC5 and F2R9_TC14) strains with their respective parent strain, a slight difference was observed (Figure 1A and Table 1). For a comparative purpose, we included an hvKpn-rmpA positive strain (K. pneumoniae 10271), and, as expected, it did not sediment well, showing a turbid supernatant (OD_{600} 0.6; Figure 1A).

### Materials and Methods

#### Plasmid \( K \)-locus determination

The \( K \)-locus was determined using Kaptive tool (Wyres et al., 2016). The plasmid profile was determined by Kaiser protocol, the size of the plasmid was determined by whole plasmid sequencing (see section “Material and Methods”). The hypermucoviscosity was determined by string test. Sedimentation assay was determined according to Bachman et al. (2015), and statistical analysis were determined.

#### Gene Products

Gene products; scrK, Fructokinase; fruA, PTS-system, IIC component.

#### Serum resistance determination

Serum resistance determination; S, sensitive; IS, intermediately sensitive. Nomenclature and abbreviations: -, Negative; +, Positive; NA, not applicable; ND, not determined; S, susceptible; R, resistant; Rif, Rifampicin; Tel, Tellurium; and NA, Nalidixic Acid.

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pKV8917 Plasmid Acquisition Increases Capsule Production and Impacts Phagocytosis by THP-1 Macrophages but Not Serum Resistance

Glucuronic acid (GA) is a type of uronic acid precursor sugar nucleotide and a common component of capsules and O antigens (Shu et al., 2009). GA was measured to determine the amount of CPS produced in the parent, cured, and transconjugant strains. Figure 1B indicates that K. variicola 8917 produced 63.2 µg of GA µg⁻¹ CFU⁻¹, whereas ΔpKV8917 showed a significant reduction in the production of GA of more than 50% (P < 0.001); however, the elimination of the plasmid did not completely abolish the capsule. Growth on MacConkey agar indicated that ΔpKV8917 and 8917 strains maintained the mucoid colony phenotype typically associated with the presence of a capsule.

Phagocytosis is a crucial process to eliminate pathogens at the early stages of infection; however, many bacteria have developed strategies to avoid phagocytosis and have become “resistant” to the action of neutrophils and macrophages, which are major components of the innate immune response (Medzhitov, 2007). Capsules are by far the main virulence factors that protect bacteria against the host immune response by inhibiting phagocytosis and lysis by complement and antimicrobial peptides (Paczosa and Mecsas, 2016). In this study, we compared the level of phagocytosis in the parent, cured, and transconjugant strains. Our results showed that K. variicola 8917 is less phagocytosed by THP-1 macrophages than the cured strain ΔpKV8917 (P < 0.001; Figure 1C). Additionally, the F2R9_TCY and F2R9_TC14 transconjugants were also less phagocytosed by macrophages compared with the parent strain F2R9 (P < 0.0001; Figure 1C). These data correlate with the changes brought by the loss or acquisition of pKV8917 over the production of CPS. This effect was not observed in E. coli J53-2 or its transconjugants because despite containing the pKV8917 plasmid, these strains did not display hmv-like properties (Figure 1A and Table 1).

Loss or Acquisition of the pKV8917 Plasmid Drives Differences in Virulence

The level of virulence was evaluated in healthy and diabetic mice in an induced sepsis model. The mortality rate of K. variicola 8917 at 6 × 10⁸ CFU was 100% after 72 h; this effect was not observed for the ΔpKV8917 strain at the same bacterial dose (Figure 2A). Otherwise, mice were infected with 4 × 10⁸ CFU K. variicola 8917 and ΔpKV8917, 40 and 80% survival rates were observed, respectively (Figure 2B). Clearly, the virulence of the ΔpKV8917 strain decreased, although the mortality rate was 40% at the highest dose inoculated. Note that lower doses of the ΔpKV8917 strain did not result in mortality; moreover, the inoculation of K. variicola F2R9, which was used as a non-hmv strain, did not result in mortality, even at high doses.

Most importantly, mice that were infected with 6 × 10⁸ of F2R9_TC14 transconjugant showed 40% mortality at 96 h (Figure 2A); however, mortality of 0% was observed at a lower dose (4 × 10⁸ CFU; Figure 2B). Thus, the level of virulence of the F2R9_TC14 transconjugants was higher than that of F2R9.

Diabetes is considered a risk factor for infections caused by K. pneumoniae, especially for hypervirulent strains (Catalan-Najera et al., 2019); however, the risk factor for hmv-like strains has not been assessed. Using a diabetic mouse model, we evaluated whether this condition could affect the survival rate when mice are challenged with an hmv-like strain compared with a non-hmv strain. Diabetic mice inoculated with 6 × 10⁸ CFU of K. variicola 8917 showed 100% mortality at 48 h in contrast to healthy mice (72 h; Figure 2C). A fast killing was not observed with ΔpKV8917 and K. variicola F2R9 in diabetic mice at the same bacterial dose inoculated (Figures 2A,C).

FIGURE 1 | (A) Mucoviscosity, (B) glucuronic acid production, and (C) phagocytosis killing assays for K. variicola 8917, ΔpKV8917, F2R9, E. coli J53-2, and transconjugants. Each symbol represents the number of biologically independent experiments. Data are presented as the mean ± standard deviation of three independent experiments. ***P < 0.0001; ****P < 0.001; and **P = 0.005. Strain 10271 corresponds to a hypervirulent/hypermucoviscous K. pneumoniae mpH⁻¹/KL2⁺ (Catalan-Najera et al., 2019) and was used as a reference for non-sedimentation behavior.
FIGURE 2 | Survival curves of K. variicola 8917, ΔpKV8917, and F2R9_TC14 in a healthy mouse model using two different CFUs (A, B). Diabetic mice were infected with $6 \times 10^8$ CFU (C) of K. variicola 8917 and ΔpKV8917. Non-hmv strain K. variicola F2R9 was inoculated at the higher dose tested ($8 \times 10^8$) and did not cause death. Significance was calculated with the Mann–Whitney two-tailed U-test.

pKV8917 Has an IncFIB Backbone but a Unique Genetic Makeup

A diagram of pKV8917 (343-kb) is shown in Figure 3. The complete sequence of the pKV8917 plasmid was compared with p15WZ-82_Vir (282,290 bp; Yang et al., 2019), pVIR_030666 (235,448 bp; Lu et al., 2018), and unnamed2 plasmid (212,079 bp) recovered from a CTX-M-15-producing K. pneumoniae strain KSB1_10J (Gorrie et al., 2018). pKV8917 exhibited 99% identity with 49% coverage to the virulence plasmid p15WZ-82_Vir and pKSB1_10J unnamed2 plasmid, both self-transmissible (Figure 3). The common regions encoded conjugation machinery (tra), tellurium resistance cluster (ter), histone-like nucleoid protein (H-NS), plasmid maintenance genes (parAB/sopAB), type I toxin-antitoxin (TA) system (hokB), and error DNA repair system (umuCD) and several hypothetical proteins (Figure 3). Another common feature was the incompatibility group IncFIB; however, pVir_030666 contained an additional replicon family, IncH1B (Lu et al., 2018).

Further sequence analysis showed the identification of regions aligned with the chromosomes of P. coffeiphila and R. terrigena NCTC13098 (Figure 3). We identified a cellulose biosynthesis cluster (bcs) and a fructose-dependent (fru) phosphotransferase system (PTS) and two genes encoding a sucrose-dependent (scr) PTS. The bcs cluster was also harbored by R. terrigena NCTC13098 and five genomes of K. variicola (VRCC00297, VRCC00299, VRCC00486, VRCC00296, and VRCC00300 from the BioProject PRJEB18814). The fru-PTS system had a significant identity to a PTS from Pantoea coffeiphila. Genes encoding the fru-PTS and the cellulose biosynthesis cluster were flanked by two insertion sequences (IS6), suggesting its mobilization and integration into the pKV8917 plasmid (Figure 3). The two genes encoding a sucrose PTS (scr) had a significant identity with genes from R. terrigena NCTC13098. The transcriptional regulators araC and iclR were also found in pKV8917 (Figure 3); they exhibited high similarity and coverage to araC and iclR regulators found in Kosakonia species. Although pKV8917 shares regions with known plasmids, its genetic structure is unique.

Moreover, we identified two clinical isolates reported as hmv-like Kv18L and TUM14115 obtained from endodontic infection and bacteremia, respectively (Harada et al., 2019; Nakamura-Silva et al., 2020). A BLAST search of the rmpADC and rmpA2 genes against these two genomes confirmed their absence. Genome alignment between 8917, Kv18L, and TUM14115 showed a ~95% nucleotide identity (Supplementary Figure 4). BLAST analysis revealed that TUM111415 lacked a plasmid replicon because all contigs corresponded to chromosomal hits, whereas Kv18L showed several plasmid-associated contigs. As there were no plasmid assembly data for Kv18L and TUM111415 that apparently lack plasmid replicon, we were not able to perform a plasmid comparison between pKV8917 and other plasmids of K. variicola strains with hmv-like properties.

Analysis of Capsule Locus of hmv-Like Klebsiella variicola

Analysis of mutations in the capsule protein-coding genes showed that although the entire K-Locus is conserved among KL114 capsule-type strains, there are several amino acid substitutions across the CPS proteins (Supplementary Figure 5 and Supplementary Table 3). The glycosyltransferase (GT_KL114) was the most diverse gene with 16 aminoacid
Diverse capsule types were identified in other hmv-like K. variicola, Kv18L and TUM14115 isolates, which were KL34 and KL125 capsule types, respectively. Additionally, manCB genes, which are responsible for GDP-D-mannose synthesis, a common component of capsules, were identified in F2R9 and 8917 capsule loci. On the other hand, the hypervirulent K. variicola isolates used for plasmid comparison (15WZ-82 and WCHKP030666) both possessed the KL16 capsule type.

DISCUSSION

Klebsiella variicola 8917 showed a hypermucoviscous-like phenotype; however, it does form a pellet upon centrifugation as non-hmv strains. K. variicola 8917 seems to behave as non-hmv when compared with K. pneumoniae 10271, a hv-hmv and KL2 strain (Catalan-Najera et al., 2019). We suggest that the sedimentation property in K. variicola 8917 depends on its polysaccharide, lipid composition, or hmv composition, which could vary between these hmv strains and may lead to non-sedimentation typically associated with hv-hmv K. pneumoniae.

When the pKV8917 plasmid was acquired by K. variicola F2R9, we noted that its transconjugants (F2R9_TC5/TC14) produced more capsule than K. variicola 8917 and F2R9 (Figure 1B). Notably, the J53-2_TC7 and J53-2_TC5 transconjugants maintained the same production level of CPS production, consistent with the results of mucoviscosity assay and the absence of hmv-like (Figures 1A, B). The F2R9_TC5 and F2R9_TC14 transconjugants were also less phagocytosed by macrophages compared with their parent strain F2R9 (P < 0.0001). However, the serum resistance assay showed a serum sensitive grade, and so it seems that hmv-like is advantageous to evade phagocytosis but do not prevent the bactericidal action of the complement system (Table 1). Phagocytosis and production of CPS correlate with the changes

FIGURE 3 | Alignment of the complete sequence of pKV8917 with two plasmids of hypervirulent K. variicola strains (p15WZ-82_Vir and pVir_030666) and one plasmid of a multidrug-resistant CTX-M-15-producing K. pneumoniae strain (pKS81_10J). We also included the chromosomes of R. terrigena NCTC13098 and P. coffeiphila. % of nucleotide identity for each region aligned is indicated in the color legend.
brought about by the loss or acquisition of pKV8917, but inconsistently, the measurement of mucoviscosity was lower than expected; although the string test was positive, the filament in transconjugants was not the same size as that produced by K. variicola 8917. A similar result was also observed in the work by Yang et al. (2019), who transferred the p15WZ-82_Vir virulence plasmid to several Klebsiella strains. Further analysis of the effects of acquiring an hmv-like-plasmid by strains with the same capsule type may explain why hmv was not exhibited at the same level as the plasmid-harboring strain.

To date, the number of reports worldwide of hmv-like K. variicola is limited (Garza-Ramos et al., 2015; Harada et al., 2019; Imai et al., 2019; Nakamura-Silva et al., 2020), and particularly in Mexico, there are no additional reports. The misclassification of K. variicola as K. pneumoniae may impact the identification of K. variicola hmv-like strains.

Altogether, mating experiments and plasmid curing suggest the presence of a novel mechanism involved in both hmv and hyperproduction of the capsule in K. variicola 8917 in an unknown interacting network. Furthermore, we assessed the impact of the acquisition of the pKV8917 plasmid in a non-hmv isolate and its elimination in K. variicola 8917 on virulence level and phagocytosis. The F2R9_TC14 and F2R9_TC5 transconjugants were less phagocytosed than the F2R9 strain. This effect is likely associated with the presence of hmv-like and hyperproduction of the capsule, which have been related to high resistance to phagocytic uptake by macrophages (Paczosa and Mecas, 2016). Although K. variicola F2R9 possesses “avirulent” behavior, F2R9_TC14 had a mortality of 40%; thus, the acquisition of pKV8917 led to an increase in virulence. In contrast, the ΔpKV8917 strain was phagocytosed in greater numbers than K. variicola 8917, which is consistent with a reduction in capsule, hmv production, and virulence.

Other plasmid-encoded virulence determinants could also contribute to virulence, including the tellurite resistance cluster (terZABCDE and terW), which is highly distributed in hvKpn strains and other species of the K. pneumoniae complex with hmv phenotype (Arena et al., 2015; Garza-Ramos et al., 2016; Martinez-Romero et al., 2018). The tellurite resistance cluster has been considered as a defense system involved in the pathogenesis (Yang et al., 2019).

Likewise, our data showed that diabetic mice challenged with hmv-like K. variicola 8917 have worse outcomes, with rapid death of the total mouse population occurring in less time and are more prone to infection than mice infected with a non-hmv strain (F2R9).

Klebsiella variicola occupies environmental niches (Rodriguez-Medina et al., 2019); thus, pKV8917 may have originated from environmental strains such as Pantoea sp., Raoultella sp. (formerly Klebsiella terrigena), and Kosakonia sp. It has been previously suggested that K. variicola could be transmitted to humans by phytosyenosis (Martinez-Romero et al., 2018; Rodriguez-Medina et al., 2019). K. variicola shares some environmental niches with Pantoea, Raoultella, and Kosakonia, and which may interchange genetic material. Seemingly, plasmid recombination events occurred among K. variicola and K. pneumoniae bacterial species resulting in a chimeric plasmid with some genetic regions belonging to Pantoea, Raoultella, and Kosakonia and incFIB self-transmissible plasmid from K. pneumoniae (Supplementary Figure 6). This scenario is plausible considering that the incompatibility group IncFIB identified in pKV8917 is highly disseminated among plasmids of clinical and community strains of K. pneumoniae (Wyres et al., 2020).

The RmpADC-pathway is a complex regulatory cascade that triggers hmv in hypervirulent K. pneumoniae. We suspect that hmv-like-associated genes in K. variicola 8917 may come from environmental strains and may be related to genes involved in metabolic fitness, particularly those implicated in sugar transport such as the fructose- and sucrose-PTS. The metabolic cost for assembling the capsule and the production of hmv could be high and requires more precursors for completing both processes; this may explain the role of the PTSs in hmv-like and capsule synthesis (Figure 3).

The biosynthesis of cellulose by Klebsiella has not been described. This feature is maintained throughout the plant kingdom and some bacterial species, such as Rhizobium sp. and Acetobacter sp. (Saxena et al., 1994). Intriguingly, we identified other isolates of K. variicola harboring the cellulose cluster. Cellulose is an extracellular complex polysaccharide; however, we do not know whether cellulose-loci in K. variicola 8917 could add to the hmv-like. Finally, the involvement of the transcriptional regulators araC and iclR in regulating vast processes such as carbon metabolism, virulence, and the glyoxylate bypass is intriguing because they may be participating indirectly in the regulatory circuit of the hmv-like (Gallegos et al., 1997; Molina-Henares et al., 2006). Additional experiments are required to assess the role of these genes over the hmv-like in K. variicola 8917.

The increasing number of hmv-like K. pneumoniae isolates is intriguing and suggests that other pathway(s) are involved. Recently, Ernst et al. (2020) reported novel capsule phenotypes mediated by deletions or single chromosomal mutations of core capsule biosynthesis genes among multidrug-resistant K. pneumoniae ST258. They found hypercapsulated mutants due to a missense mutation in the wzc gene (G565S). This substitution was searched in the K. variicola 8917 isolate; however, no mutation in the wzc gene was found in that position, and we did not detect mutations in the rcsAB or lon genes (data not shown). Comparing the KL114 capsule-type of three different strains, we identified capsule locus variation throughout the cps. Noteworthy, the cps loci of both K. variicola were more similar than the KL114 of K. pneumoniae (Supplementary Figure 5). KL114 is a rare capsule type among K. pneumoniae; the presence of subtypes within KL114 may not have been addressed due to limited sequence data. Nonetheless, the composition and diversity of K. variicola capsule loci deserve to be studied in detail.

Hypermucoviscosity is detected more frequently, and multiple pathways can trigger it; the implications of this phenotype when a hypervirulent background is absent deserve further analysis because this phenotype may play a role as a virulence factor. In conclusion, the acquisition of pKV8917 plasmid confers the hmv-like phenotype triggering an increase in capsule and virulence, especially with risk in diabetic populations. This study proposes
a novel plasmid-borne mechanism that could be disseminated among the *K. pneumoniae* complex.

**DATA AVAILABILITY STATEMENT**

The sequences of the chromosome and plasmid of *K. variicola* 8917 have been deposited in GenBank under accession numbers: CP063403 and CP063404, respectively. Bioproject PRJEB7831.

**ETHICS STATEMENT**

The animal study was reviewed and approved by Biosafety Committee at National Institute of Public Health.

**AUTHOR CONTRIBUTIONS**

UG-R and NR-M conceived and designed the experiments. NR-M, MD, MA, and HV-T performed the experiments. UG-R, NR-M, and LL-A provided analytical tools. UG-R, JS-S, VA, and JM-B contributed reagents and materials. NR-M, EM-R, and UG-R wrote the manuscript. All authors contributed to the article and approved the submitted version.

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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.2020.579612/full#supplementary-material

**Supplementary Figure 1** | ERIC profiles of the (a) 8917 and F2R9 parent strains and transconjugants F2R9_TCS14 and JS3-2_TCS7 and (b) ΔpKV8917 cured strain. The red symbol represents a possible plasmid fragment of ~500 bp.

**Supplementary Figure 2** | Phylogenetic tree based on the concatenated sequence of pgi, leuS, and pyrG genes. The phylogeny was generated with the clustering method neighbor joining using MEGA6 software.

**Supplementary Figure 3** | Plasmid profile based on alkaline lysis. Lines 1 and 10, *E. coli* NCTC 50192: plasmids were used as molecular size markers (154-, 46-, and 7-kb; Bachman et al., 2015); lines 2-3, *K. variicola* 8917 (pKV8917); line 4, ΔpKV8917 (cured strain); lines 5-6, JS3-2_TCS5 and JS3-2_TCS7; lines 7-8, F2R9_TCS5 and F2R9_TCS14; and line 9, F2R9.

**Supplementary Figure 4** | Genome sequence alignment of 8917 vs TUM14115 and Kv1BL obtained from *K. variicola* isolates.

**Supplementary Figure 5** | Phylogeny of isolates with capsule-type KL114. The phylogenetic tree was based on the concatenated amino acid sequence of 19 proteins that make up KL114 capsule type. *K. variicola* strains 13450 and 8917 are indicated with a triangle, *K. pneumoniae* strain QMP with black square and black circle is showed. In addition, the comparison of the complete KL114 capsule locus. CDSSs are presented as arrows. Gray bars indicate regions of similarity, and darker regions indicate a higher degree of similarity. Red arrows correspond to mannose synthesis and processing proteins (ManC and ManB). The number of mutations found across the cps is represented with blue bars. Detailed information on the sites of mutation is in Supplementary Table 3.

**Supplementary Table 3**

**Supplementary Figure 6** | A model for how the pKV8917 may have originated. (A) *K. variicola* is ubiquitous in soil, plants, animals and water. These natural niches possess a diverse bacterial population and allow *K. variicola* to coexist with other endophytes and bacteria of animal carriage. *Parotepa* sp. *Kosakonia* sp. and *Raoultella* sp. were donors of genetics regions to *K. variicola* 8917 possibly by horizontal transfer. Raoultella sp. is isolated more frequently from animal gut and sewage so that *K. variicola* 8917 might have acquired the cellulose biosynthesis cluster (clos) in this spot. Our hypothesis is that hypermucoviscosity was acquired by bacterial species from the environment. By phytobiontes, humans become infected by *K. variicola* 8917 which could be transported to the community (B) or hospital setting (C) in this scenario *K. variicola* coexist with *K. pneumoniae*. IncFIB plasmids (IncFIBp) are usually associated with multidrug-resistant and virulent strains; IncFIBp carries the transference module (tra) and tellurium resistance operon (ter). IncFIBp was transferred by means of conjugation to *hmv*-like *K. variicola* strain. IncFIBp serves as the backbone for the incorporation of the genetic elements acquired from environmental strains resulting in the structure pKV8917; pKV8917 could be transfer the *hmv*-like to species of the *K. pneumoniae* complex (D) but also acquire virulence and multidrug-resistance genes.
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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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