Supplemental Figures

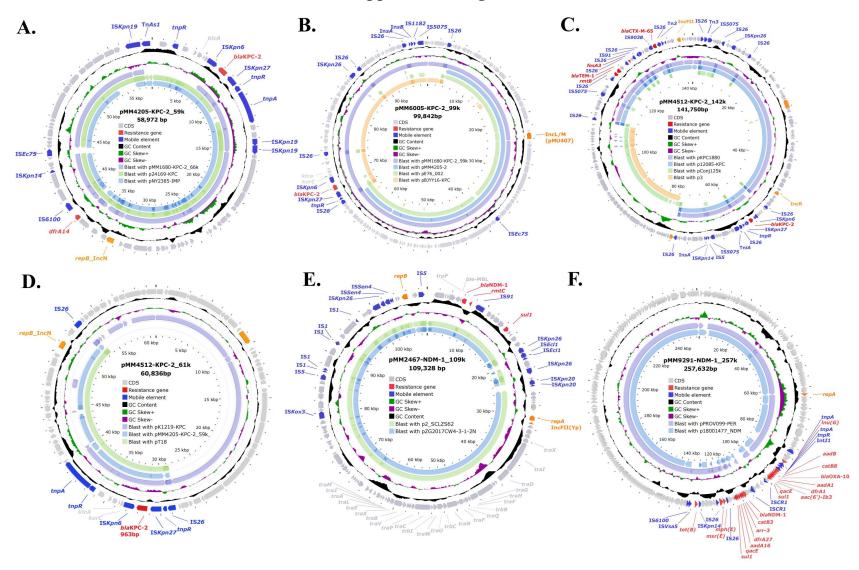


Fig. S1. Circular sequence alignment of plasmids bearing blakpc-2 or blandm-1. The plasmids originated from two sources: our study and other similar plasmids available from the NCBI database through BLAST. (A) Both pMM4205-KPC-2 59k (58,972 bp) and pMM1680-KPC-2 66k (66,924 bp), were highly similar to p24169-KPC (MN891676, 69k) (100% identity and 96% coverage) from clinical Klebsiella pneumoniae, as well as to pNY2385-IMP (CP096922, 62k) (100% identity and 93% coverage) from Citrobacter freundii. (B) The pMM6005-KPC-2 99k (99,842 bp), a large recombinant plasmid, showed significant similarity to pMM1680-KPC-2 99k based on the IncL/M plasmid with IS26-mediated mobile sequences. The MM4512 strain hosted double bla_{KPC-2} in two separate plasmids, one of which was pMM4512-KPC-2 142k (141,750 bp) (C), classified as an IncFII-R recombinant plasmid. This complex plasmid contains a variety of mobile elements and several resistance genes, including bla_{KPC-2}, bla_{CTX-M-65}, bla_{TEM-1B}, fosA3 and rmtB. It was similar to pKPC1880 (CP061347, 169k) (100% identity and 74% coverage) and p12058-KPC (MN842292, 142k) (100% identity and 73% coverage) from K. pneumoniae. A type IV secretion system (T4SS), similar to p3 (CP132686, 35k) from K. pneumoniae and pConj125k (MK033499, 125k) from *E. coli*, was inserted by IS26. Additionally, pMM4512-KPC-2 61k (60836 bp) (**D**), carrying bla_{KPC-2} variant (963 bp) with bla_{KPC-2} 1 (AY034847, VCV70871) (93% identity and 99% coverage), showed the highest similarity to pK1219-KPC (CP137007, 140k) (100% identity and 76% coverage) from Citrobacter koseri and pMM4205-KPC-2 59k (IncN) in our study. (E) The pMM2467-NDM-1 109k (109,328 bp) was highly similar to p2 SCLZS62 (CP082170, 111k) (100% identity and 97% coverage) from *Raoultella planticola* and pZG2017CW4-3-1-2N (CP065352, 108k) (100% identity and 95% coverage) from *K. pneumoniae*. **(F)** The pMM9291-NDM-1_257k (257,632 bp), a recombined plasmid, was similar to pPROV099-PER (CP120536, 259k) (100% identity and 91% coverage) and p18001477_NDM (CP098041, 273k) (100% identity and 91% coverage) from *Providencia rettgeri*.

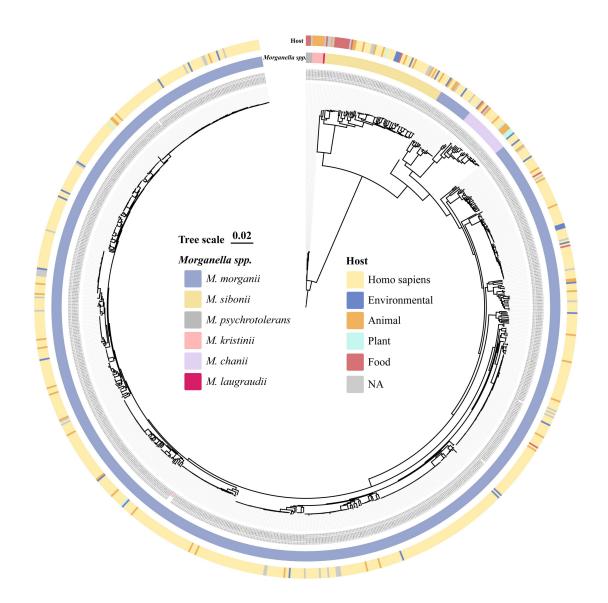


Fig. S2. Phylogenetic tree of global 968 Morganell spp. isolates. Employing the Fastani identification method, a genomic relatedness threshold of ANI (Average Nucleotide Identity) greater than 95% was established to delineate homologous species. This criterion successfully classified all 968 Morganella spp. into the 6 species. Reference Sequences: Morganella morganii (GCA_006094455.1), Morganella sibonii (GCA_039099315.1), Morganella chanii (GCA_000633515.1), Morganella kristinii (GCA_001676055.1), Morganella laugraudii (GCA_003996855.1), Morganella psychrotolerans (GCA_001676155.1).

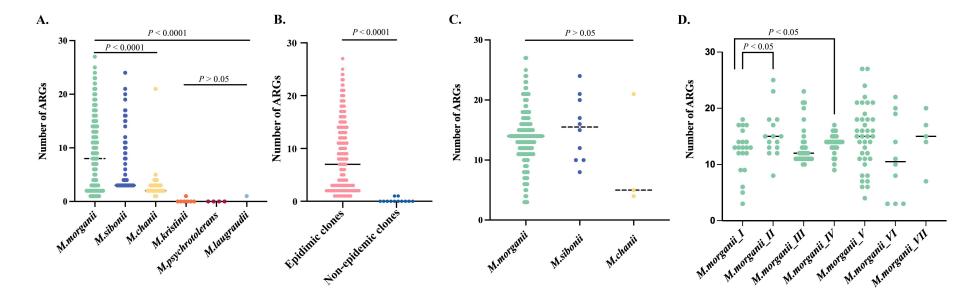


Fig. S3. Comparative analysis of ARGs quantities among different *Morganella* spp. (A) Among 968 *Morganella* spp. isolates, the number of ARGs in epidimic clones (*M. morganii*, *M. sibonii* and *M. chanii*) was greater than in non-epidemic clones (*M. kristinii*, *M.psychrotolerans* and *M. laugraudii*), as determined by ANOVA and two sample *t*-test (B). (C) Three types of CRMs producing KPC or NDM did not exhibit significant differences in the number of ARGs. (D) Among 177 *M. morganii* bearing *bla*_{KPC} or *bla*_{NDM} genes, there were differences in the number of resistance genes among some phylogroups.