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The Environmental Exposures and Inner- and Intercity Traffic Flows of the Metro System May Contribute to the Skin Microbiome and Resistome

Highlights

- Intraday time is the primary determinant of metro microbiome and resistome composition

- Human commensals and clinical ARGs are of higher relative abundance in p.m. samples

- Line-specific environmental exposure also modulates the microbial community

- Potential cross-boundary ARG transmission event was captured

Authors

Kang Kang, Yueqiong Ni, Jun Li, ...,
Morten Otto Alexander Sommer,
Christopher John Webster,
Gianni Panagiotou

Correspondence

gianni.panagiotou@hki-jena.de

In Brief

Kang et al. present a metagenomic analysis of the Hong Kong metro system. They show a reliable and dynamic view of the diurnal flux of microbial transmission and recurrence affected by the traffic flow and establish a baseline for metagenomic studies examining human interactions with public transit.
The Environmental Exposures and Inner- and Intercity Traffic Flows of the Metro System May Contribute to the Skin Microbiome and Resistome

Kang Kang, Yueqiong Ni, Jun Li, Lejla Imamovic, Chinmoy Sarkar, Marie Danielle Kobler, Yoshitaro Heshiki, Tingting Zheng, Sarika Kumari, Jane Ching Yan Wong, Anand Archna, Cheong Wai Martin Wong, Caroline Dingle, Seth Denizen, David Michael Baker, Morten Otto Alexander Sommer, Christopher John Webster, and Gianni Panagiotou

1Systems Biology and Bioinformatics Group, School of Biological Sciences, Faculty of Sciences, The University of Hong Kong, Hong Kong S.A.R., China
2Novo Nordisk Foundation Center for Biosustainability, Technical University of Denmark, 2800 Kgs. Lyngby, Denmark
3Healthy High Density Cities Lab, HKUrbanLab, The University of Hong Kong, Hong Kong S.A.R., China
4School of Public Health, The University of Hong Kong, Hong Kong S.A.R., China
5School of Biological Sciences, Faculty of Science, The University of Hong Kong, Hong Kong S.A.R., China
6Department of Earth Sciences, The University of Hong Kong, Hong Kong S.A.R., China
7Division of Landscape Architecture, The University of Hong Kong, Hong Kong S.A.R., China
8Systems Biology & Bioinformatics Unit, Leibniz Institute for Natural Product Research and Infection Biology, Hans Knöll Institute, Jena, Germany
9Lead Contact
*Correspondence: gianni.panagiotou@hki-jena.de
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SUMMARY

The skin functions as the primary interface between the human body and the external environment. To understand how the microbiome varies within urban mass transit and influences the skin microbiota, we profiled the human palm microbiome after contact with handrails within the Hong Kong Mass Transit Railway (MTR) system. Intraday sampling time was identified as the primary determinant of the variation and recurrence of the community composition, whereas human-associated species and clinically important antibiotic resistance genes (ARGs) were captured as p.m. signatures. Line-specific signatures were notably correlated with line-specific environmental exposures and city characteristics. The sole cross-border line appeared as an outlier in most analyses and showed high relative abundance and a significant intraday increment of clinically important ARGs (24.1%), suggesting potential cross-border ARG transmission, especially for tetracycline and vancomycin resistance. Our study provides an important reference for future public health strategies to mitigate intracity and cross-border pathogen and ARG transmission.

INTRODUCTION

The skin is not merely a physical barrier between the body and the environment but functions as a complex contact interface for a rich and influential microbial community. Indeed, the human skin and its microbial residents play an essential role in health and disease. The human skin microbiome has been shown to have a unique signature and to maintain stable characteristics at the strain level over time (Oh et al., 2016). Despite this resistance to colonization and disruption, the skin is a source of potential infection, especially in vulnerable individuals. For instance, various Propionibacterium acnes strains were identified as potential determinants for acne, the most common skin condition (Fitz-Gibbon et al., 2013). Skin from over 90% of atopic dermatitis (AD) patients is highly colonized with the pathogen Staphylococcus aureus (Leyden et al., 1974), whereas Staphylococcus epidermidis was found to modulate the host immune response to inhibit S. aureus (Cogen et al., 2010a, 2010b). The microbial composition of the human skin can be affected by our lifestyle choices, including cohabitants, dwelling environment, and even our pets (Lax et al., 2014). In high-density megacities, mass transit systems also serve as platforms for the exchange of microbes between millions of humans and the built environment (Hong Kong’s Mass Transit Railway [MTR] has a daily ridership of around 4.7 million, for example; MTR Corporation, 2017), induced by the high mobility of the population and constant interaction through skin-surface contact. Mass transit systems therefore represent ideal targets for studying the composition and dynamics of urban microbial communities. They constitute passive samplers with unique niches and a potentially higher propensity for microbial accumulation, exposure, and transmission, created through large concentrations of people from diverse urban backgrounds interacting in relatively confined spaces (Be et al., 2015).

To characterize the urban metro system microbiota, several 16S rRNA-based studies have been conducted in high-density cities, including New York and Hong Kong (Leung et al., 2014; Robertson et al., 2013). A recent shotgun metagenomic study performed in the New York metro system characterized the
citywide urban microbial geographic distribution using samples collected from station surfaces (Afshinnekoo et al., 2015). This study reported potential health risks associated with the microbial communities found in the stations, including the presence of pathogens and antibiotic resistance as well as constant daily recurrence in the vast bacterial ecology. A similar study of the Boston metro system (Massachusetts Bay Transportation Authority [MBTA]) highlighted that different surface types and materials are occupied by different microbes with high variation in functional capacity and pathogenic potential (Hsu et al., 2016). Recently, more studies of metro and urban microbiomes have started under the collaborative framework of the Metagenomics and Metadesign of the Subways and Urban Biomes (MetaSUB) International Consortium (MetaSUB International Consortium, 2016). Previous metagenomic studies of metro systems were conducted by sampling directly from surfaces or indoor air masses. However, to obtain a more focused picture of how microbial communities occupying the surfaces in a metro system may affect human health, studies focusing on the contact interface—by sampling the skin microbiome—are of significant interest. Ultimately, a key objective of performing microbial profiling in a transit system is to identify possible health-related risks, as evidence has accumulated of the transmission of clinically significant disease from surface to hand (Bhalla et al., 2004; Weber et al., 2010) as well as community-acquired antibiotic-resistant infections (Ho et al., 2007; Spellberg et al., 2008; Vandenesch et al., 2003). Therefore, characterizing shifts in the microbial consortia an individual carries after using mass transit is a more direct investigation than intensive sampling of accumulations and communities on built surfaces (Gibbons et al., 2015; Prescott et al., 2017; Zimmerman et al., 2014). Furthermore, because mass transit is characterized by intermittent passenger loads (fluctuating over peak and off-peak hours) and inherent tidal effects between downtown and uptown regions, time-series-based studies with distinct sampling times throughout the day and repeated sampling days offer a more reliable and dynamic view of the diurnal flux of microbial transmission and the propensity of recurrence induced by the variation in traffic flow. Finally, investigating the correlations between microbial composition and the underlying geographical or topological characteristics of different metro lines as well as their connectivity to the underlying urban matrix (street networks, buildings, and public spaces) is crucial because these connections govern human mobility and interactions in high-density environments and, hence, influence the rate of microbial contact and transmission (Adams et al., 2015). Several interesting questions that may inform public health surveillance and urban planning were raised. What is the composition of the microbiome and its potential health risk (e.g., antibiotic resistance and pathogenicity) we may acquire when exposed to mass transit? Will the microbiome inherit tidal effects of the traffic flow and vary over traveling time? How might urban density and morphology influence the microbial composition and health risk it carries? To what extent is a city’s unique metro microbiome influenced by connecting routes to a neighboring metropolitan region or country? To answer these questions, we conducted a metagenomic study of the Hong Kong MTR system (Figure 1A) with a cross-border rail connection to the neighboring large city of Shenzhen in mainland China. Shotgun metagenome sequencing followed by taxonomic and functional analysis was performed on samples collected from the surface of human palms after contact with metro railcar handrails (Figure 1B). Our sampling covered different metro lines and was conducted at different time points across the day with weekly replicates. To determine whether the microbiome and resistome vary over the day and differ by distinct metro lines, we looked for time-specific and line-specific signature species.
and antibiotic resistance genes (ARGs), respectively, and then we attempted to track potential transmission events of ARGs. Further, to reveal how urban design could influence microbial ecology and biosafety issues, we investigated the correlations among metro network topology, ecological features (e.g., community diversity), and health risk levels, such as clinically important ARG abundance, ARG dissemination potential, and pathogenic potential. Our study provides a useful framework for obtaining a comprehensive view of mass transit systems as modulators of human microbial contacts, informing public health planning regarding monitoring and health risk management.

RESULTS

Microbial Communities Acquired by MTR Exposure
Taxonomic profiles of the relative abundance were generated at each taxonomic level for each sample (Data S1A and S1B). Most bacterial reads were derived from four major bacterial phyla: Actinobacteria, 51.3% ± 8.5%; Proteobacteria, 26.6% ± 9.3%; Firmicutes, 11.4% ± 3.0%; Bacteroidetes, 2.3% ± 0.7%; and 8.4% ± 1.0% for others (Figure 2A). P. acnes was the dominant organism, accounting for 29.1% ± 7.0% of the community (Figure 2B). In the 10 most populated species, 8 were common human and skin commensals, including P. acnes, Micrococcus luteus, Propionibacterium humerusii, Acinetobacter baumannii, S. epidermidis, Escherichia coli, and S. aureus. The yeast Malassezia globosa (0.72%), Enterobacteria phages (0.28%), and Propionibacterium phages (0.13% in 27 subtypes, including 8 abundant subtypes) are the most abundant non-bacterial organisms. Not surprisingly, the population of Propionibacterium phages is positively correlated with the abundance of Propionibacterium (Pearson’s correlation coefficient (r) = 0.72, p = 6.7e−9) and 11 Propionibacterium species, including P. acnes (r = 0.71, q = 1.7e−6). All 8 types of abundant Propionibacterium phages co-occurred with their hosts at both genera and species levels. To achieve sufficient resolution for analyses of phage-host interactions and pathogenicity, strain-level taxonomic construction was performed for two major species, P. acnes and S. epidermidis, with well-documented reference strains using a reference-based approach (Oh et al., 2014). In the 89 strains quantified as P. acnes, C1 was identified as the leading strain (making up 12.6% ± 3.2% of the species) in 44 of the 48 samples, followed by PA2 (6.4% ± 0.6%) and P.acn17 (5.5% ± 1.8%) (Figure S1A; Data S1C). Notably, we observed that different Propionibacterium phages showed drastic variability in their predilection to multiple P. acnes strains. In general, the overall abundance of Propionibacterium phages is correlated with 24 P. acnes strains (r = 0.47–0.72, q < 0.1). Specifically, phage ATCC29399B_C shows correlation with 36 strains, followed by PHL037M02, P1.1 and P100_1, correlated with 15, 14, and
Microbial Communities Significantly Differ by Sampling Time

Sampling time (a.m. versus p.m.) was identified as the major determinant of the shifts and recurrences in community composition and diversity. Simpson’s alpha-diversity indices notably decreased from a.m. to p.m. (0.90 to 0.85, paired t test, p = 1.6e-4; Table S1), accompanied by a consistent increase in P. acnes abundance (25.7% to 32.5%, paired t test, p = 2.1e-4; Figures 2B and 2C). The ER line, the only cross-border line connecting to Shenzhen in mainland China, appears as the only outlier. At the species level, the abundances of 140 species significantly decreased from a.m. to p.m. (paired t test, p < 0.05), with a median rank of their relative abundance in all analyzed species of 419.5/744 (Figure 3B). We defined these species as a.m. signature species. In only 48 species, the population expanded from a.m. to p.m., including Enterobacteria phages and Propionibacterium phages in 5 subtypes, with a median rank of 232/744 (Figure 3B). These species were defined as p.m. signature species. The Matthew effect can be used as an analogy for delineating the dynamics of microbial communities “the rich (a.m.) get richer (p.m.) and the poor get poorer” in their abundance (Merton, 1968). By analyzing the pathogenicity of the time-specific signature species, only 3 potential opportunistic pathogenic species of low abundance (<0.3%) were a.m. signatures, whereas 11 were p.m. signature species, led by P. acnes and S. mitis. The total abundance of potential opportunistic pathogenic species is also dramatically increased (paired t test, p = 5.9e-6) but becomes insignificant when excluding P. acnes (paired t test, p = 0.25; Figure 3A). To determine the driver(s) in the niche expansion of P. acnes, we performed comparisons at the individual strain level. SK137 and hdn-1 significantly increased from a.m. to p.m., whereas HL035PA1 decreased (Wilcoxon signed-rank test on ranks, p < 0.05). Notably, none of these altered strains have been previously reported as pathogenic strains. When assigning the species to ecological categories (including different habitats and physiological traits; Data S3A) using the PATRIC database (Wattam et al., 2014), we detected a significant increase p.m. for the categories of skin-associated, disease and/or human disease-related, hosted and/or human-hosted, non-motile, and salinity-tolerant species and a decrease in marine species (as well as in soil, aquatic, plant-hosted, and motile species when removing the outlier ER line) (paired t test, p < 0.05; Figure 3A). Interestingly, for some non-human-associated species with high abundance, we also observed a high fluctuation in abundance regardless of sampling time points. Such examples include the fourth most abundant species, Burkholderia, a nitrogen-fixing soil organism, with a relative abundance of 1.97% ± 3.94%, Bradyrhizobium elkanii (19th, 0.48% ± 1.64%), and Labrenzia alexandrii (20th, 0.48% ± 0.96%). Burkholderia and L. alexandrii showed remarkable variance in abundance among the three sampling days (ANOVA test, p < 0.05, Burkholderia: 0.00%, 0.86%, and 5.04%; L. alexandrii: 0.91%, 0.53%, and 0.00% for the 3 days). B. elkanii had extremely high abundance only in day 1 p.m. samples in the ER and MOS lines (10.47% and 4.53%; the relative numbers were 1.07% and 1.31% in a.m. compared with 0.13% as the mean for other samples). The PATRIC categories of animal-hosted species, aquatic species, marine species, non-antimicrobial resistance (AMR) species, thermo-tolerant species, and species with multiple altitude preferences (1,000 m, 10 m, and 1 m) showed instability among the 3 sampling days, and none of these categories are human-related. One possible explanation is that this variation could be driven by non-human factors, such as humidity and aerosols, because weather conditions during the sampling days differed (Table S1). Based on this evidence, we speculate that daily traffic flow is the driving factor in shaping the microbial communities of a public transit system. Human-associated species are universally more abundant than non-human-associated species, and they tend to become more abundant under high traffic density during the day; conversely, species with low relative abundance are unlikely to be human commensals, and they are further outcompeted by the abundant human-associated species during high-traffic hours before...
recovering their ecological niche during non-traffic hours. Naturally, the abundant species show higher inter-day stability in their composition, whereas, conversely, the less abundant non-commensal species show greater stochastic variations among different sampling days.

**Microbial Communities Are Line Dependent**
In addition to sampling time, community diversities (Table S1) and dissimilarities revealed that the microbial compositions were also highly influenced by the metro line features (Figure 2D). The distances among communities from different lines become smaller from a.m. to p.m., except for the ER line, the only cross-border line linked to mainland China, which is also the only interchangeable line of the most isolated line in the MTR system, the MOS line. The coordinates of the a.m. and p.m. communities from the MOS line locate closely to the p.m. and a.m. communities from the ER line, respectively. The coordinate swapping may reflect the effect of the traffic tide induced by the unique interchange. IndVal tests (Dufrene and Legendre, 1997) were performed to identify line-specific signature species: species that are relatively abundant only in one line compared with all other lines. Line-specific signatures were investigated among all samples or among only a.m. or p.m. samples. In general, more a.m. signature species (88) could be detected than p.m. signatures (59) or all-day signatures (78) (Figure 3C) because the community dissimilarities dramatically decreased p.m. The most isolated line, MOS, showed a markedly greater number of signature species (55) both a.m. and p.m. as well as the highest community diversity a.m. (one-against-all t test, p = 8.0e−5; Figure 3D). The ER and KT lines were also characterized by 20 and 16 a.m. signature species, respectively, before losing their discriminatory power p.m. The stability of the signature species is also line-dependent: only 21 of the 88 a.m. signature species could retain specificity p.m., 17 of which are MOS line signatures; only 4 species are signatures for the other 7 lines. When assigning species into PATRIC categories, line-specific signature categories could be observed (one-against-all Kruskal-Wallis test on ranks, p < 0.05; Figure 3D). The MOS and ER lines reported the highest abundance of soil species and species with AMR in all samples but motile species only p.m. The MOS
line, which is an entirely aboveground ground line running alongside the Shing Mun Channel, a polluted brackish river, is the most populated line for aquatic species a.m. and sewage species both a.m. and p.m. The WR line, which passes through a mountain range in the New Territories, showed the highest abundance of the species with a habitat preference for altitude around 1,000 m, whereas a low abundance of these environmental categories was observed in underground urban lines such as ISLAND, TW, and TKO. Species that were originally isolated in Hong Kong were most abundant in the TW line, an urban line with the highest traffic volume (Table S2), which connects the densely populated areas in Kowloon with the Central station on Hong Kong Island. All of these observations regarding microbial composition are associated with line-specific environmental exposure. To estimate interline transmissions and track potential sources of the heterogeneously mixed p.m. communities, we employed the Bayesian SourceTracker algorithm (Knights et al., 2011) to estimate the contributions of the a.m. communities, assuming that the p.m. communities (sink communities) originated, transmitted, and were mixed from 8 independent a.m. communities (source communities), and each line a.m. community is contributing to a particular proportion of the p.m. communities. Not surprisingly, microbial relocation between the ER and MOS lines was detected at a high level. The a.m. community of the ER line contributes a notable proportion to multiple p.m. communities, including the MOS, TC, WR, and TW lines (Figure S2). The two lines with the longest interchange routes to the ER line, the TKO and ISLAND lines, are relatively less colonized by the a.m. community of the ER line. As the most isolated line, the MOS line contributes a very limited percentage to all non-ER lines, supporting its capacity to retain the specificity of signature species p.m.

**The MTR Resistome and Functionome**

The unique characteristics of the ER line observed in the above analysis triggered our interest in the possibility of cross-border transmission of ARGs via the MTR system. Intensive analysis of the resistome profile in the MTR lines was performed to explore the ARG mechanism (Data S3B), gene family (Gibson et al., 2015; Data S3C), previously annotated clinical relevance (Munck et al., 2015), and dissemination potential (STAR Methods). In general, antibiotic efflux (469.2 ± 182.6 RPM [reads per million reads]) was revealed as the most abundant characterized mechanism in the MTR communities, followed by target protection (146.3 ± 26.7) and inactivation (137.3 ± 35.3), target alteration (69.1 ± 13.5), and replacement (26.7 ± 9.8). Overall, we identified 136 ARG families (the overall abundance was 322.6 ± 62.1 RPM), including 24 clinically important ARG families (14.1 ± 5.1 RPM) (Data S3C). The most abundant ARG families were fluoroquinolone-resistant DNA topoiso-merase (42.8 ± 6.6 RPMK), *aph(3’)* double prime (34.5 ± 5.1), RND antibiotic efflux pump (13.4 ± 5.8), ABC antibiotic efflux pump (13.2 ± 3.0), and *macB* (9.2 ± 2.9). The most abundant clinically important ARG families were *ermC* (3.6 ± 2.7 RPMK), *msrA* (3.3 ± 2.0), *tetK* (1.7 ± 0.7), *ant4(4’)-Ia* (1.5 ± 0.6), and *tetM* (1.0 ± 0.4). The abundance of clinically important ARG families showed higher inter-sample variability than non-clinical families, suggesting a potentially higher effect of human activities. Therefore, a systematic analysis of the time- and line-specific signature ARG families was a logical follow-up.

**The p.m. Signature of the ARG Families Is Highly Associated with Human Activities and Clinical Antibiotic Practice**

When comparing the resistome between a.m. and p.m. communities, no differentiation was captured in the abundance of the ARG mappable reads. Similarly, the abundance of clinically important ARGs and community-wide dissemination potential increased modestly without statistical significance (Figure 3A). The community-wide dissemination potential was estimated from the horizontal gene transfer (HGT) rate of each ARG family (Data S4; STAR Methods). At the ARG family level, the abundance of 4 ARG families significantly decreased from a.m. to p.m. (*ant(2’), ant(3’), aac(6’)-Ib*, and SubclassB2; all non-clinical), whereas 9 families increased, including clinically important families like *tetM, ermC, vanB*, tetracycline resistance ribosomal protection protein (tetRPPP), *tetA-B*, *tetM-tetW-tetO-tetS* (tetMWOS), *vanD*, and *mpf* (paired t test, p < 0.05; Figure 3A). Resistance against the clinically widely used antibiotics tetracycline, vancomycin, erythromycin, and mexitilllin was highly observed in p.m. communities. These observations led us to pose a new question: is the increase in clinically relevant antibiotic resistance contributed only by the colonization process of human-associated species, which occurs equally in all lines, or is it also accompanied by inter-line ARG dispersion carried by the transmission of line-specific signature species? To answer this question, we investigated the line-specific ARG dispersion and taxonomic contributors.

**The ER Line’s Resistome Signature Suggests Potential Cross-Border Antibiotic Resistance Transmission**

Besides sampling time, we also examined the specific characteristics of each line and observed that the MOS line has a significantly higher abundance of all ARG reads. The highest abundance of clinically important ARGs and highest community-wide dissemination potential were observed in the TC and ER lines (one-against-all Kruskal-Wallis test on ranks, p < 0.05; Figure 3D). The ER and RC lines showed a high abundance of the ARG family *bial* beta-lactamase repressor gene as the major contributor to the high dissemination potential. The MOS line is ranked third in terms of community-wide dissemination potential, driven by the high HGT rate of the *aph(3’)* family. These 3 lines are uptown lines with predominantly aboveground tracks. Similar to the line-specific signature species, the ER and MOS lines also lead in terms of number of signature ARG families: 5 for the ER line, 4 for the MOS line, and only one for each of the other lines (emrA for the TC line) (Figure 3C). The MOS line has more non-clinical signature ARG families (*ant(3’), SubclassB3, and GES*) and only one clinically relevant family (*vanC*); in contrast, all ER line signature ARG families are related with clinically used antibiotics (Figure 3D), and 4 of 5 (*tetA, tetO, tetRPPP, and tetMWOS*) are involved in resistance against tetracycline, besides one against vancomycin (*vanC*). Among these 5 families, 4 (excluding tetMWOS) are a.m. signatures without specificities p.m.

**Busy Lines Do Not Show a Higher Abundance of Clinically Important ARGs or Higher ARG Dissemination Potential**

To disclose whether civic characteristics such as urban density and metro network topology are also influencing the
microbial diversity, line signatures, resistome composition, and its transmission potential, we performed an additional association analysis using a hybrid urban movement network operationalized by integrating the MTR topological network (Figure 4A; Data S5A), the adjoining street network linked to MTR station entrances (Figure 4B; Figure S3; Data S5B), the traffic volume (The Transport and Housing Bureau of the Government of the Hong Kong Special Administrative Region, 2014; Table S2), as well as the distances to the Central station and Shenzhen (mainland China) (Data S5A). The distance to Central station was used to distinguish downtown and uptown lines, whereas the distance to Shenzhen was introduced to capture intercity transmission events. We correlated these features with community-wide and health-related observations, including community diversity, ARG abundance, clinically important ARG abundance, and community-wide ARG dissemination potential. At the community scale, diversity (in Simpson’s index) was observed to be positively correlated with the line’s average distances to Central station (Figures 4C and 4D; significant p.m., Pearson’s correlation coefficient $r = 0.41$, $p = 0.046$). The diversity differences between p.m. and a.m. (Simpson’s index p.m.–a.m.) were positively correlated with the line average edge betweenness ($r = 0.47$, $p = 0.019$; Figures 4C and 4D). These findings indicate that isolated or uptown lines tend to have higher community diversities and greater decreases in the community diversities during the day. In relation to the resistome, strong negative correlations were captured between the clinically important ARG abundance and the mean betweenness of the road connectivity networks ($r = 0.42$ to $0.37$, $p < 0.05$; Figures 4C and 4D), whereas a negative correlation was also observed for the community-wide ARG dissemination potential in the a.m. samples ($r = 0.48$ to $0.41$, $p < 0.05$; Figures 4C and 4D). There was also a strong positive correlation between the increase of dissemination potential (p.m.–a.m.) and the distance to Central station ($r = 0.44$, $p = 0.033$; Figure 4C). Based on these correlations, the isolated uptown lines tend to have a higher abundance in clinically important ARGs and ARGs with higher dissemination potential. The distance to Shenzhen (mainland China) was negatively correlated with the overall...
Correlations for three individual ER line signature ARG families have higher ARG input during the day. We also detected the values), suggesting that lines closer to Shenzhen tend to as its increment (weak correlation calculated from line mean

correlated with the distance to Shenzhen, especially in a.m. (tetA was positively corre-
lated with distance (r = 0.50, p = 0.012; Figures 4 Ca n d 4 D ), The increased abundance of tetA that could be gained over a day, which is consistent with our previous observations: an ER line a.m.

Correlations between Species, ARG Families, and Biosynthetic Gene Clusters

to further characterize the functionality of the microbial communities in connection to the observed resistome profile, we quantified and compared the abundance of the biosynthetic gene clusters (BGCs) (Donia et al., 2014; Data S3D), which has also been suggested by the MetaSUB Consortium as a standard component for urban microbiome characterization (MetaSUB International Consortium, 2016). Among the 22 retrieved BGC types, 3 types were identified as p.m. signature types: ladderane, bacteriocin, and lantipeptide. Bacteriocin and lantipeptide are BGC types closely related to antibiotic resistance. Six types were identified as a.m. signature types: type I polyketide (t1pks), trans-AT polyketide (transatpks), type III polyketide (t3pks), terpene, indole, and other types of polyketide (otherks) (Figure 3A). Among the 7 line-specific signature BGC types, 4 were associated with the WR line (Figure 3D). SparCC correlations were calculated among BGC types and ARG families (Data S6). The p.m. signature lantipeptide was detected to be highly associated with multiple ARG families (Figure 6C). The BGC types aryl polycyline (arylpolyene) and homoserine lactone (hserlactone) were also highly associated with ARG families. Investigating the ARG co-occurring and co-excluding networks (Figures 6A and 6B), we could observe competing modules in both a.m. and p.m. networks: one human or clinic-associated module with the msr, ble, mpr, erm, ctx-m, and tet families and one non-clinical module with the ade, bae, tol, sox, cfr, md, abc, and ant families. In the a.m. network, stronger co-occurring links could be observed in the non-clinical module, whereas the connections among the clinical families were relatively weak. In contrast, stronger links among clinical links could be identified in the p.m. network, whereas the non-clinical module was losing internal connections. These observations also support our conclusions from the time-specific ARG analysis.

Co-occurring relationships and taxonomic links were observed between the abundances of ARG families, BGC types,
and their contributing species (Figure 6C; Data S7). At the phylum level, the most abundant phylum, Actinobacteria, contributed to both ARGs and BGCs. Proteobacteria and Firmicutes contributed more to ARGs but less to BGCs, whereas Bacteroidetes showed links only to ARGs. The dominant species, P. acnes, was not determined to be a major contributor to any ARG family. S. aureus, E. coli, S. suis, K. oxytoca, S. arlettae, and Williamsia sp. D3 were the top-ranking species contributing to the clinically important ARG families. Notably, we could observe a high abundance of S. aureus in the TC, MOS, and ER lines. The ER line was the most populated line for S. suis, K. oxytoca, and S. arlettae, whereas the MOS line showed a higher abundance of Williamsia spp. The abundance of the top clinical ARG contributing species was always led by clinically important ARG families. Notably, we could observe a high abundance of S. aureus in the TC, MOS, and ER lines. The ER line was the most populated line for S. suis, K. oxytoca, and S. arlettae, whereas the MOS line showed a higher abundance of Williamsia spp. The abundance of the top clinical ARG contributing species was always led by clinically important ARG-enriched lines (the ER and TC lines). Investigating the line signature ARG families, S. suis was the major contributor to the ER signature families tetA and tetO. The ER line was also the most abundant line for S. suis, blal, an ARG family with an extremely high HGT rate, had a distinctly higher abundance in the TC and ER lines, and its only contributing species, S. aureus, was also abundant in the TC (second) and ER (fourth) lines. For the MOS leading family aph(3'), also with a high HGT rate, multiple contributors (S. maltophilia, A. baumannii, and P. aeruginosa) were identified, and the MOS line was validated to have a high abundance of A. baumannii, followed by S. maltophilia. On the other hand, BGC-type contributing organisms were more extensively distributed throughout the whole community (4,303 taxa contributing to 40 BGC types) in comparison with the ARG contributors (only 2,116 taxa linked with 136 ARG families). The greatest BGC contribution was from M. luteus, followed by P. acnes, Gordonia bronchialis, Kytococcus sedentarius, and Corynebacterium jeikeium. For the 4 ARG-associated BGC types mentioned above, P. acnes was the dominant contributor to lantipeptide, whereas bacteriocin was evenly linked with multiple species from Streptococcus, Kocuria, Burkholderia, Haemophilus, and other genera, and aryl polyene and homoserine lactone shared A. baumannii as their leading contributing species.

**DISCUSSION**

Previous studies have investigated the microbial composition of mass transit aerosols within the New York City and Hong Kong MTR (Leung et al., 2014; Robertson et al., 2013) as well as station and train surfaces in New York City and Boston (Afshinnekoo et al., 2015; Hsu et al., 2016). The skin microbial community should also be appreciated as a pivotal component of such indoor investigations. The skin microbiota is adapted to the human body and may influence human health through its capacity to travel with individuals, shed from the skin, adhere to indoor
surfaces, and recolonize other individuals (Lax et al., 2014). Our study focuses on the characterization of skin microbiota acquired when exposed to a public mass transit environment such as the Hong Kong MTR. Like other studies of train surfaces, station surfaces, and aerosols (Afshinnekoo et al., 2015; Hsu et al., 2016; Leung et al., 2014; Robertson et al., 2013), the metro microbial communities recovered from palm samples in the Hong Kong MTR were largely derived from human skin commensals regardless of the sampling sites. In addition, shotgun sequencing provided significantly higher resolution of the taxonomic profiles and identified *P. acnes* as the dominant species, which was not identified in early 16S rRNA sequencing-based studies, leading to a potential bias in the interpretation of microbial communities (Leung et al., 2014; Robertson et al., 2013). By applying shotgun sequencing, we could delve into the strain-level analysis, which identified putative pathogenicity, interactions with distinct phage subtypes, and strain-specific variation and contribution to morning versus afternoon conditions.

Regarding the resistome, the resistance genes against chloramphenicol, tetracycline, and beta-lactamase reported by previous studies (Afshinnekoo et al., 2015; Hsu et al., 2016) were also observed in our samples. Our investigation of the skin microbial community enabled the precise characterization of the antibiotic resistance, based on the relatively high stability of the microbial communities (Oh et al., 2016). In comparison with the Boston metro metagenomic study (Hsu et al., 2016), more resistance to widely used clinical antibiotics, including vancomycin, erythromycin, and mexitilin, could be detected in our samples with daily recurrence. Besides the clinically important ARGs, this study also investigated and estimated the ARG dissemination potential at the family level and community scale. Routine screening for clinically important ARGs and assessment of their dissemination potential would allow improved detection of potential health risks induced by antibiotic resistance outbreaks, geographical transmission, or cross-species migration.

The universal stability of the skin microbiome (Oh et al., 2016) promotes the reliability and the reproducibility of longitudinal analyses across the sampling intervals used in our study: short intervals as in different sampling times in a day and median intervals as in different sampling days in successive weeks. The sampling intervals have successfully assisted with exploring the effect on community diversity and stability from the intraday tidal effect as well as the inter-day variation. Intraday sampling time was identified as the major driver of the variation and recurrence of microbial composition and diversity in conjunction with the ARGs and BGCs carried by the community composition. When evaluating the inter-day stability, the less abundant environmental species (not human-hosted organisms) showed significantly higher instability, further suggesting their potential competitive disadvantages under the pressure of human traffic and the influence of non-human factors, such as indoor climate (Brodie et al., 2007; Kelley and Gilbert, 2013). Taxonomic contributors to the ARG families were also captured for the time- and line-specific signature ARGs, with strong co-occurring trends, serving as evidence of the presence of signature ARGs and their transmission events (Hiramaatsu et al., 2014; Lowy, 2003). Tracing ARG-contributing species could serve as a starting point for strategies that might inhibit ARG transmission and dissemination in the future. Furthermore, in our analyses, we considered the taxonomy, resistome, and BGC profiles as interrelated components. Taxonomic contributors and associated BGC types were regarded as direct and implied evidence for the divergence observed in the resistome (Doria et al., 2014).

When exploring the resistome, clinically important ARG families display higher variances, suggesting greater influence from human traffic. Notably, most p.m. signature ARG families are closely affiliated with clinical use of antibiotics. The specificity of line-specific signatures decreased from a.m. to p.m., in accordance with a previous evaluation of the stability of site-specific signatures of skin microbes (Oh et al., 2016). By reviewing the time- and line-specific ARG analyses in conjunction, we captured some evidence for intra- and intercity ARG transmission. As a notable case, we captured 4 tetracycline- and 2 vancomycin-resistant ARG families as universal p.m. signature families: tetMWOS, tetRRPP, tetM, tetA-B, vanB, and vanD. All of these families were identical or highly associated with the ER line signature families. The ER line showed a markedly higher abundance in these families a.m., and then these families gained in abundance in all lines and became universal p.m. signature families, whereas the ER line lost specificity in these families (Figure 3A). From these findings, one may hypothesize that the ER line, the only cross-border line linked to mainland China, may serve as a potential source of clinically important ARGs, especially against tetracycline, which is a commonly used antibiotic in China’s swine feedlots (Wu et al., 2010) and can be detected in the soil in the Pearl River Delta area (Li et al., 2011), where the cities of Hong Kong and Shenzhen are located. It appears that these ARGs disperse into different lines in the MTR system with the carriage of human traffic and finally become relatively abundant in every line in the later hours of the day.

Furthermore, from our study, robust patterns in microbial community dynamics and clinically relevant ARG transmission were observed. To evaluate the potential real health effect of such transmission events, the population size of the microbial community becomes a key factor. In our preliminary study, we also collected samples by swabbing the train surfaces, following the same procedure as used in the Boston study (Hsu et al., 2016). Our DNA amounts from both palm and surface samples (<1 ng/μL) were orders of magnitude lower than those in the original study (40–220 ng/μL) (Hsu et al., 2016), which suggests a relatively lower overall biomass of the MTR microbiome and relatively higher safety when using the MTR, as the transmission of pathogens and ARGs is kept at a low level. The low DNA recovered may be attributed to the effectiveness of the antibacterial nano-silver-titanium dioxide coating (NSTDC) applied in the MTR (MTR Corporation, 2006). Assuming that no such antibacterial practice was applied, the robust patterns of the clinically relevant ARG transmission could be amplified, resulting in increased health effects. We propose that public policy makers should continue to pursue and evaluate evidence of the effect of antimicrobial strategies applied to public transit systems and checkpoints (e.g., border control regions, airports and aircraft, cross-border trains and buses, etc.), especially between regions with different norms and regulations in industrial and clinical antibiotic usage. In addition to maintaining and expanding the silver nanoparticle coating prophylaxis, public restroom
sanitation could be improved, and hand sanitizer dispensers could be provided at the exits of metro, train train, and bike-sharing stations, airports, and hospitals to reduce the transmission of pathogens and clinically relevant ARGs. At a higher level, the growing understanding of the contribution of microbial communities to human health should be reflected in design of built environments and materials and equipment for public use with attention to the microbiome. Studies have found that architecture and ventilation design affect community diversity (Kembel et al., 2014). Furthermore, given the effect of substrate on community composition, future work on the dynamics of optimizing spaces and surfaces for human health in relation to microbiota is needed.

In summary, our investigation of the microbiome associated with public transit reveal that communities are decidedly shaped by human interactions and traffic flows, and we hope to have established a baseline for further studies by incorporating multidimensional influences from the host, time, environment, geography, and varied civic characteristics. In future studies, more sampling time points could be considered to delineate more exquisite intraday dynamics, especially the time points before and after morning and evening peaks, respectively. A dynamic and surveillance of the microbial communities responsive to and shaped by the public transit systems could serve as a reference for public policy-making regarding the prevention of ARG spread via intercity and cross-border traffic and protect millions of citizens from potential health hazards in daily life.

STAR METHODS

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SUPPLEMENTAL INFORMATION

Supplemental Information includes three figures, two tables, and seven data files and can be found with this article online at https://doi.org/10.1016/j.celrep.2018.06.109.

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AUTHOR CONTRIBUTIONS


DECLARATION OF INTERESTS

The authors declare no competing interests.

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REFERENCES


### STAR METHODS

#### KEY RESOURCES TABLE

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### CONTACT FOR REAGENT AND RESOURCE SHARING

Further information and requests for data and resources should be directed to and will be fulfilled by the Lead Contact, Gianni Panagiotou (gianni.panagiotou@hki-jena.de).
EXPERIMENTAL MODEL AND SUBJECT DETAILS

Sample collection
Each sample was a pooled community from the palms of 6 healthy individuals (without symptoms of skin disease or other illness) who maintained contact with different handrails in the Hong Kong MTR system for 30 minutes (Figures 1A and 1B). For each of the 8 urban lines in the Hong Kong MTR system, samples were collected in the morning (9:00 to 11:30 am) and evening (19:00 to 21:30 pm). Each line was sampled 3 times within 3 consecutive weeks. For each sample, we collected metadata recording the line, sampling date and time, sampling route interval and weather condition of the sampling day (Table S1).

In each sampling section, volunteers followed a standard procedure: 1. Wash hands using normal soap and distilled water for 90 s before boarding the train; 2. Touch different handrail surfaces in a running train of a specific line for approximately 30 minutes; 3. Swab each palm for 2 minutes using the ESwab Collection & Transport System (BD company, pre-moisturized with 0.15M NaCl and 0.1% Tween 20). To provide adequate DNA yield for library preparation of each sample, one swab was used for all 12 palms. Swabs were stored in 15 mL LabServ tubes on ice and transported to a -20°C freezer within an hour. DNA extractions were performed within 24 hours after sample collection.

An extra 6 negative controls were prepared (3 from the same palms without touching any surface for 30 minutes, 3 from unused swabs). The DNA yields were undetectable by Qubit (< 0.007 ng/μl in 50 μL elution buffer) for all negative controls, suggesting that the DNA that we collected originated from the MTR train surfaces but not from the swabs, buffer solutions or remaining bacteria on palms after washing.

Ethics approval
All participants in the sampling procedure were researchers from the University of Hong Kong, and gave written informed consent. Sequencing reads from human genome were removed before analyses. The study was approved by the Human Research Ethics Committee (HREC) of the University of Hong Kong.

METHOD DETAILS

DNA extraction and metagenomic sequencing
Microbial DNA was extracted with the PowerSoil DNA Isolation kit (MO BIO Laboratories) following the official protocol. 0.95 ± 0.41 ng DNA was obtained for each sample. The extracted DNA was used to construct shotgun metagenomic libraries using the Nextera XT kit following the standard protocol, which is optimized for the library construction for low DNA input (1 ng). The metagenome sequencing was performed with Illumina HiSeq 1500 (101bp PE) at the Centre for Genomic Sciences of The University of Hong Kong. The sequenced reads were processed for quality control to remove the adaptor regions, low quality reads and the reads from human genome following the previously described steps (bwa mapping against GRCh37/hg19 reference genome using mem algorithm, extracting reads with > 95% identity) (Li et al., 2016). 75% to 94% of the reads passed the quality control, with a mean value of 90.4%. The human reads occupied 39% to 78% of the clean reads, with a mean value of 43.8%.

The total DNA amount obtained (0.95 ± 0.41 ng) was much lower than in other studies which collected samples by swabbing surfaces in metro stations directly (Afshinnekoo et al., 2015; Hsu et al., 2016), but comparable to previous human skin metagenomic projects (Oh et al., 2014). Besides the disparity in the microbial density between the handrail and the colonized palm, another possible explanation for the low DNA collected could be the effect of the antibacterial Nano-Silver-Titanium Dioxide Coating (NSTDC) applied to MTR surfaces (MTR Corporation, 2006). The DNA library construction was optimized for the low DNA input and shotgun metagenomic sequencing was performed for all samples. We used our previously developed method to perform functional metagenomics analysis after the metagenomics sequencing (Forsberg et al., 2012; Munck et al., 2015; Sommer et al., 2009; van der Helm et al., 2017), however, the DNA amount of these samples were lower than the minimal requirements for the construction of functional libraries.

Taxonomic profiling
From the metagenomic sequencing data, a million reads were randomly sub-sampled from each sample for taxonomic profiling. These reads were mapped to the NCBI non-redundant (nr) database by DIAMOND (Buchfink et al., 2015) using the default settings. To estimate the taxonomic composition of the samples, the Lowest Common Ancestor (LCA) algorithm was implemented with LCA mapper from mtools, MEGAN5 (Huson et al., 2011) (-f Detect -ms 50 -me 0.01 -tp 50). The relative abundance for each taxonomic level was distillate following the previously described pipeline (Heshiki et al., 2017; Huson et al., 2011). The 744 most abundant species (composing 95% of the total reads in all samples) were used in further statistical analyses (Data S1B).

Phylogeny
Synthetic tree of phylogeny and taxonomy for the most abundant bacterial species (composing 90% of the total reads) was acquired from OpenTree7.0 (Hinchliff et al., 2015), integrating the major studies on the phylogenetic trees for bacteria (Lang et al., 2013; Segata et al., 2013).
The abundances of ARG families were calculated as RPKM (Reads per Kilobase per Million mapped reads) (Data S3C). The overall reads were removed from the ARG mapping results. The coverage rate of each ARG was calculated by using the mappable reads from the nr database using DIAMOND blastp. If ARGs did not remain the best hits (sorted by e value) in the nr database mapping step, the DIAMOND blastp with stringent parameters (-e 1e-10–id 95, best hits reserved). The mappable reads were further mapped against the extended ResFams database (Gibson et al., 2015). The 27 sequences of clinically important ARGs described previously (Munck et al., 2013) (Data S3B), as described previously (Heshiki et al., 2017). For the ARG family information, all reads were mapped against the extended ResFams database (Gibson et al., 2015). The ARG families may have overlaps due to the family ranges defined by ResFams. In both annotation procedures, reads were mapped using DIAMOND blastp with stringent parameters (-e 1e-10–id 95, best hits reserved). The mappable reads were further mapped against the nr database using DIAMOND blastp. If ARGs did not remain the best hits (sorted by e value) in the nr database mapping step, the reads were removed from the ARG mapping results. The coverage rate of each ARG was calculated by using the mappable reads from all samples. Among 136 quantified ARG families, 98 families have a coverage > 50%, 91 families > 70% and 84 families > 90%.

Reference-based strain mapping
Strain profiling of P. acnes and S. epidermidis was conducted according to the previously suggested reference-based approach (Oh et al., 2014). The non-core region strain quantifications were used in statistical comparisons (Data S1C and S1D).

Pan-genome analysis
Pan-genome profiling was conducted by PanPhlan using default parameters for the 10 most abundant species and 10 most ARG contributing species (Scholz et al., 2016). Presence/absence and abundance profiles were generated for gene clusters.

Species categorization
The bacterial species were queried against PATRIC database (Wattam et al., 2014) for categorization. The features recorded in genome metadata include: habitat, host, mobility, salinity, altitude/depth, temperature range, oxygen requirement, pathogenicity and antimicrobial resistance. To link the PATRIC features with the taxonomic profiling at the species level, all features in any strain were aggregated and assigned to the corresponding species (Data S3A). For example, 6 strains of Vibrio ordalii were marked as aquatic strains in PATRIC, so the abundance of V. ordalii at the species level was counted for the category of aquatic species.

Identification of pathogens and potential pathogenic species
To identify opportunistic pathogens, two lists of potential pathogens (Forsberg et al., 2014; Kembel et al., 2012) were combined (Data S2A). The PATRIC database (Wattam et al., 2014) was used as a reference for taxonomic structures and species/strain names. The organisms appearing on the list were marked as opportunistic pathogens, and the species containing at least one opportunistic pathogenic strain were treated as potential opportunistic pathogenic species (Data S2B).

Identification and quantification of ARG mechanism and families
To identify ARGs and assign them to different mechanisms and ARG families, two different procedures were performed. For the ARG mechanism annotation, the one million reads subset used for taxonomic profiling were mapped against CARD database (McArthur et al., 2013) (Data S3B), as described previously (Heshiki et al., 2017). For the ARG family information, all reads were mapped against the extended ResFams database (Gibson et al., 2015). The 27 sequences of clinically important ARGs described previously (Munck et al., 2015) were also included in the extended database, if the sequenced were absent in ResFams (blastp identity < 95%). New clinically important ARG families were built for the absent ARGs in ResFams: all existing ResFams sequences were mapped against those ARGs, and the mappable genes (blastp, -e 1e-5–id 95) were classified into regarding clinically important ARG families. ARG families may have overlaps due to the family ranges defined by ResFams. In both annotation procedures, reads were mapped using DIAMOND blastp with stringent parameters (-e 1e-10–id 95, best hits reserved). The mappable reads were further mapped against the nr database using DIAMOND blastp. If ARGs did not remain the best hits (sorted by e value) in the nr database mapping step, the reads were removed from the ARG mapping results. The coverage rate of each ARG was calculated by using the mappable reads from all samples. Among 136 quantified ARG families, 98 families have a coverage > 50%, 91 families > 70% and 84 families > 90%. The abundances of ARG families were calculated as RPKM (Reads per Kilobase per Million mapped reads) (Data S3C). The overall abundances of all ARG families were calculated as RPM (Reads per Million mapped reads) (Data S4B).

Identification of virulence factors
All sequencing reads were mapped against VFDB (Chen et al., 2016), following the same procedure and parameter of ARG family annotation. The abundances of the virulence factors of the opportunistic pathogens were summarized in RPKM (Data S2C).

Community diversity and dissimilarity
To estimate the alpha-diversity of the microbial community for each sample, the Simpson diversity index was calculated with VEGAN (Dixon, 2003) based on the species-level relative abundance (Table S1).

To estimate the community dissimilarities, weighted UniFrac distances were calculated by phyloseq (McMurdie and Holmes, 2013) based on the relative abundance of species, as well as functional classes (for example, ARG family profiles). Ordinations were calculated by nonmetric multidimensional scaling (NMDS) for the illustration.

Dissemination potential estimation
To estimate the plasmid-mediated horizontal gene transfer (HGT) rate, all ARG sequences were mapped against the NCBI plasmid RefSeq database (Pruitt et al., 2014) using blastp (-e 1e-5–id 95). The protein sequences from all functional ARGs were mapped to the gene families that were acquired from the HGTtree database (Jeong et al., 2016). The HGTtree gene family of an ARG was defined by the gene family with the highest proportion of valid hits (-e 1e-5 and coverage > 50% in query or subject) in HGTtree families. RANGER-DTL (Bansal et al., 2012) was used for phylogenetic reconciliation analysis based on the gene tree and species tree provided by the HGTtree. The HGT rate in each family was calculated by dividing the number of HGT events by the total phylogenetic tree length of that family. Finally, the HGT rates were assigned to ResFams ARG families with unique mapping (Data S4A). The ARG families with extremely high HGT rate (> 100) are biaA and aph(3')-I.
The community-wide dissemination potential (Data S4B) was estimated for each sample, by as the following equation:

$$\sum \text{HGT}_i \cdot \text{RPKM}_i$$  
(Equation 1)

where:
- $i$ is an HGTree gene family;
- HGT$_i$ is the HGT rate of the family $i$;
- RPKM$_i$ is the abundance of the HGTree family $i$ in RPKM.

**Identification and quantification of biosynthesis gene clusters**

BGCs annotation was based on antiSMASH 3.0 (Weber et al., 2015) and modified as previously described (Donia et al., 2014; Ni et al., 2016). All reads were mapped against the database using DIAMOND with the same parameters as ARGs. The abundances of BGCs/BGC types were calculated as RPKM (Data S3D).

**Identification of taxonomic links for ARGs and BGCs**

For all the reads that could be mapped to ARGs/BGCs, taxonomic profiling was carried out using the same pipeline used for the 1 million read subsamples, with more stringent filtering criteria (bootstrap value > 50 in LCA mapping at species level). Therefore, the major contributing species for each ARG family or BGC type could be identified (Dataset S7).

**Morphometric modeling of hybrid rail-road urban network**

To test the hypothesized correlations between microbial transmission, loads and diversity and underlying urban morphology and connectivity, a series of network models were conducted. As a first step, initial assessment of network topology of the 8 urban lines was conducted by Cytoscape 3.3.0 (Shannon et al., 2003). Averaged edge features (EdgeBetweenness) and node features (including BetweennessCentrality, ClosenessCentrality, Degree, NeighborhoodConnectivity, NumberOfEdges, Radially) were calculated for each line (Data S5A). Line civic features (number of stations/interchange stations, number of interchangeable lines, and average station distance to Shenzhen) were also incorporated in the topology analysis.

The second step involved more detailed network modeling employing spatial Design Network Analysis (sDNA) (Cooper and Alain, 2015), a state-of-the-art urban network analysis technique developed by teams at Cardiff University and The University of Hong Kong to model multiple scale urban morphological metrics (morphometrics). The MTR network database was first coupled with the street network database by digitizing the network of subways linking platforms to station entry-exit points to adjacent street link in ArcGIS 13 and the hybrid network was cleaned, transcribed into an access graph model and subjected to sDNA analysis. Among a suite of various network morphological metrics (morphometrics) generated, betweenness centrality or through-movement potential was employed in the present study. It is the simulated counts of movement through each link in the network, given its position in the network and the geometrical and topological connectivity with other links within the network. sDNA betweenness of $x$ in a graph of $N$ links may be defined as:

$$Bt Wl(x) = \sum_{y \in R_y} \sum_{z \in \text{links}} L(y) L(z) P(z) \text{OD}(y, z, x)$$  
(Equation 2)

where:
- $y$ and $z$ are the geodesic end points;
- $R_y$ is the set of links within a defined radius from $y$;
- $L(y)$ and $L(z)$ are length of links $y$ and $z$ respectively;
- $P_z$ is the proportion of link $z$ within the defined radius.

OD is a function defined as:

$$\text{OD} = \begin{cases} 
1, & \text{if } x \text{ is on the geodesics from } y \text{ to } z \\
\frac{1}{2}, & \text{if } x \equiv y \neq z \\
\frac{1}{2}, & \text{if } x \equiv z \neq y \\
\frac{1}{2}, & \text{if } x \equiv z \equiv y \\
0, & \text{otherwise}
\end{cases}$$  
(Equation 3)

It was modeled at micro-meso-macro-level urban scales (100, 200, 300, 400, 500, 600, 700, 800, 900, 1000, 2000, 3000, 5000, 7500, 10000, 15000 m street catchments) to capture potential effects of scale on human mobility and microbial transmission (Data S5B). For each MTR station, betweenness values were aggregated at three levels: with respect to all the street links closest to all exits of a
metro station, all street links within a 50 m and 100 m catchment radii. For each MTR line, the street connectivity index was represented by the mean betweenness for all the links for the line as well as sDNA betweenness values of all stations on it. Other urban measures employed in the study included network proximity to two transportation hubs, namely Hong Kong Central station and Shenzhen Railway Station and Luohu metro link.

**QUANTIFICATION AND STATISTICAL ANALYSIS**

**Statistical analysis**

All statistical analyses were performed in R. For comparative analysis between AM and PM samples (including taxa, species categories, ARG families, BGC types, etc.), paired t tests (for abundance/proportion) or Wilcoxon signed-rank tests (for rankings) were performed. For comparative analysis among different lines or different sampling days, ANOVA (for abundance) or Kruskal-Wallis tests (for rankings) were performed. For line-specific species categories, Wilcoxon tests with one-against-all comparison were performed. To discover line-specific signatures for ARG families and BGC types, IndVal tests (Dufrene and Legendre, 1997) were performed. Alpha value of 0.05 was used for all statistical tests.

Co-occurring and co-excluding relationships among taxa, ARG families and BGC types were identified by SparCC algorithm (Friedman and Alm, 2012). Correlations with coefficient absolute value greater than 0.3 were reserved (Data S6).

Pearson’s correlation coefficients were calculated between topological or geographical features and biological features (overall ARG abundance, clinically important ARG family abundance, community-wide dissemination potential, ER line signature ARG families’ abundance, etc.). Correlations with p value < 0.05, or adjusted q value < 0.1 with Bonferroni correction for multiple comparisons, were marked as significant correlations.

**Data visualization**

R, python and corresponding packages including ggplot2, gplots, heatmap.plus and matplotlib were used for illustration of statistical results. iTOL (Letunic and Bork, 2016) was used for the phylogeny visualization. Cytoscape 3.3.0 (Shannon et al., 2003) was used to visualize the analyses incorporating network and topology.

**DATA AND SOFTWARE AVAILABILITY**

**Data from the other studies**

The sequencing data of other environmental samples used for comparison were obtained from NCBI Sequence Read Archive (accession number: ERP013563, ERP013562, ERP012177, SRP033730 and SRP061803).

**Availability of data and materials**

The sequencing reads files were deposited into the Sequence Read Archive (SRA) of the National Center for Biotechnology Information (NCBI) with accession number SRP119528 under project PRJNA413474. The datasets supporting the conclusions of this study are included within the article and the additional files.