



Yılın Ses Getiren Makaleleri:

Nozokomiyal İnfeksiyonlar

Dr. Şiran Keske
27 Aralık 2022





Innovation for infection prevention and control—revisiting Pasteur's vision

Gabriel Birgand, Raheelah Ahmad, Andre N H Bulabula, Sanjeev Singh, Gonzalo Bearman, Enrique Castro Sánchez, Alison Holmes

Lancet 2022; 400: 2250–60

Louis Pasteur has long been heralded as one of the fathers of microbiology and immunology. Less known is Pasteur's

“the dust in the atmosphere contains microorganisms which develop and multiply”

“the most putrescible liquids remain unaltered if, after heating them, they are left protected from the air, and therefore from these microorganisms”



1875

Paris maternity hospital

Lohusa humması.

64 ölüm

“Il me vient cette idée que le corps étranger quand il amène le pus, ce qui n'est pas constant, doit apporter un germe, lequel germe serait cause de la formation de pus.”

“Yabancı cisimler iltihabi akıntıya yol açıyorsa beraberinde “germ” da getiriyor olmalılar ve bu “germ” bu akıntının sebebi olmalı.

Bu olaydan sonra el hijyeninin önemi arttı.
Pasteur kendisini “el hijyeni fanatığı” olarak tanımlamıştır.



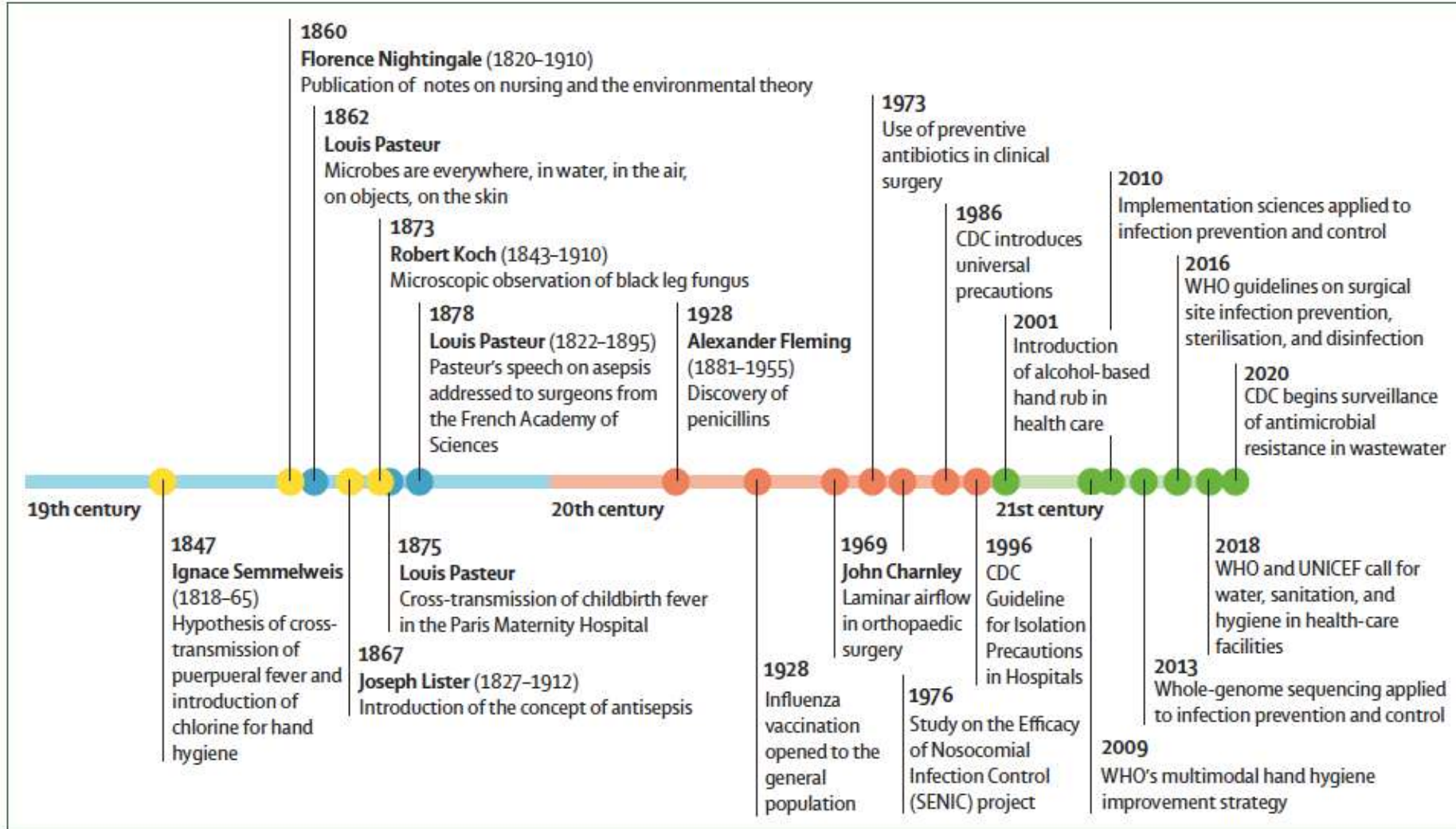


Figure 2: Timeline of selected innovations related to Pasteur's vision

“Permettez-moi de vous révéler le secret qui m’a conduit à atteindre mon but. Ma force repose uniquement sur ma ténacité.”

“Size hedefime götüren sırrı söyleyeyim. Gücüm sadece azmimde yatıyor.”
Louis Pasteur

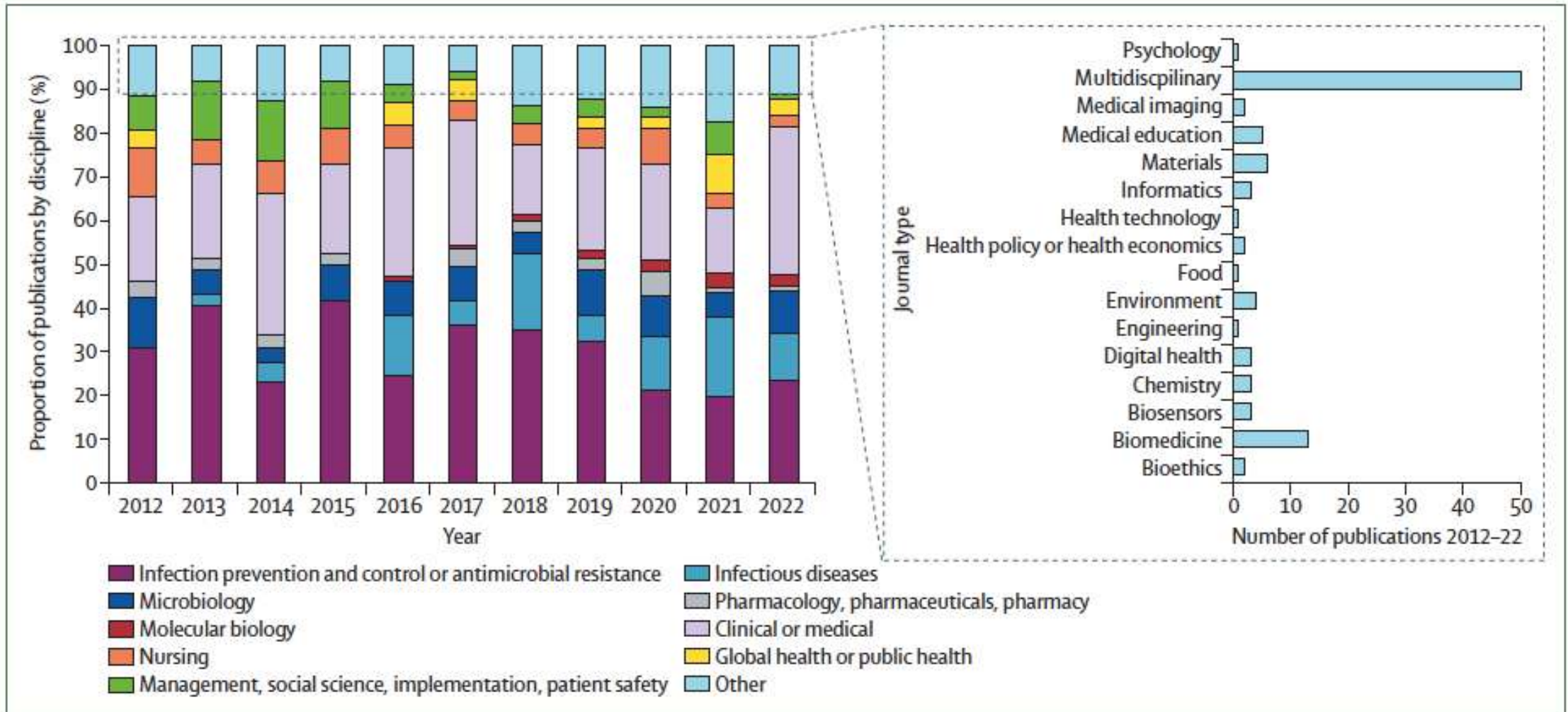


Figure 3: Dissemination of innovation development and implementation

The data show a positive trajectory to wider clinical speciality audiences but restricted learning in management and social science outputs.

inovasyon nedir?



Global burden of bacterial antimicrobial resistance in 2019: a systematic analysis

*Antimicrobial Resistance Collaborators**

Summary

Background Antimicrobial resistance (AMR) poses a major threat to human health around the world. Previous publications have estimated the effect of AMR on incidence, deaths, hospital length of stay, and health-care costs for specific pathogen–drug combinations in select locations. To our knowledge, this study presents the most comprehensive estimates of AMR burden to date.



Lancet 2022; 399: 629–55

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[https://doi.org/10.1016/S0140-6736\(21\)02724-0](https://doi.org/10.1016/S0140-6736(21)02724-0)

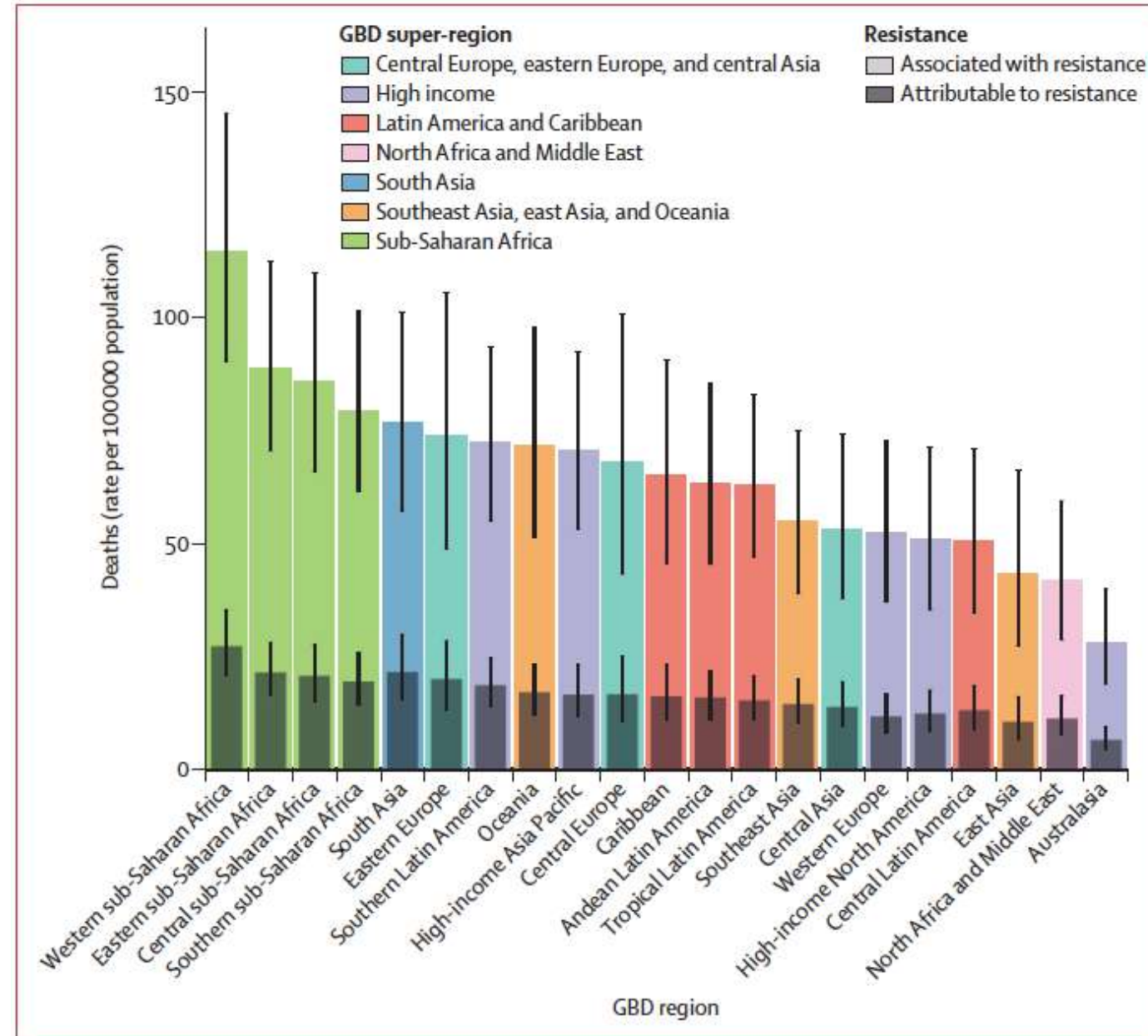


Figure 2: All-age rate of deaths attributable to and associated with bacterial antimicrobial resistance by GBD region, 2019

Estimates were aggregated across drugs, accounting for the co-occurrence of resistance to multiple drugs. Error bars show 95% uncertainty intervals. GBD=Global Burden of Diseases, Injuries, and Risk Factors Study.

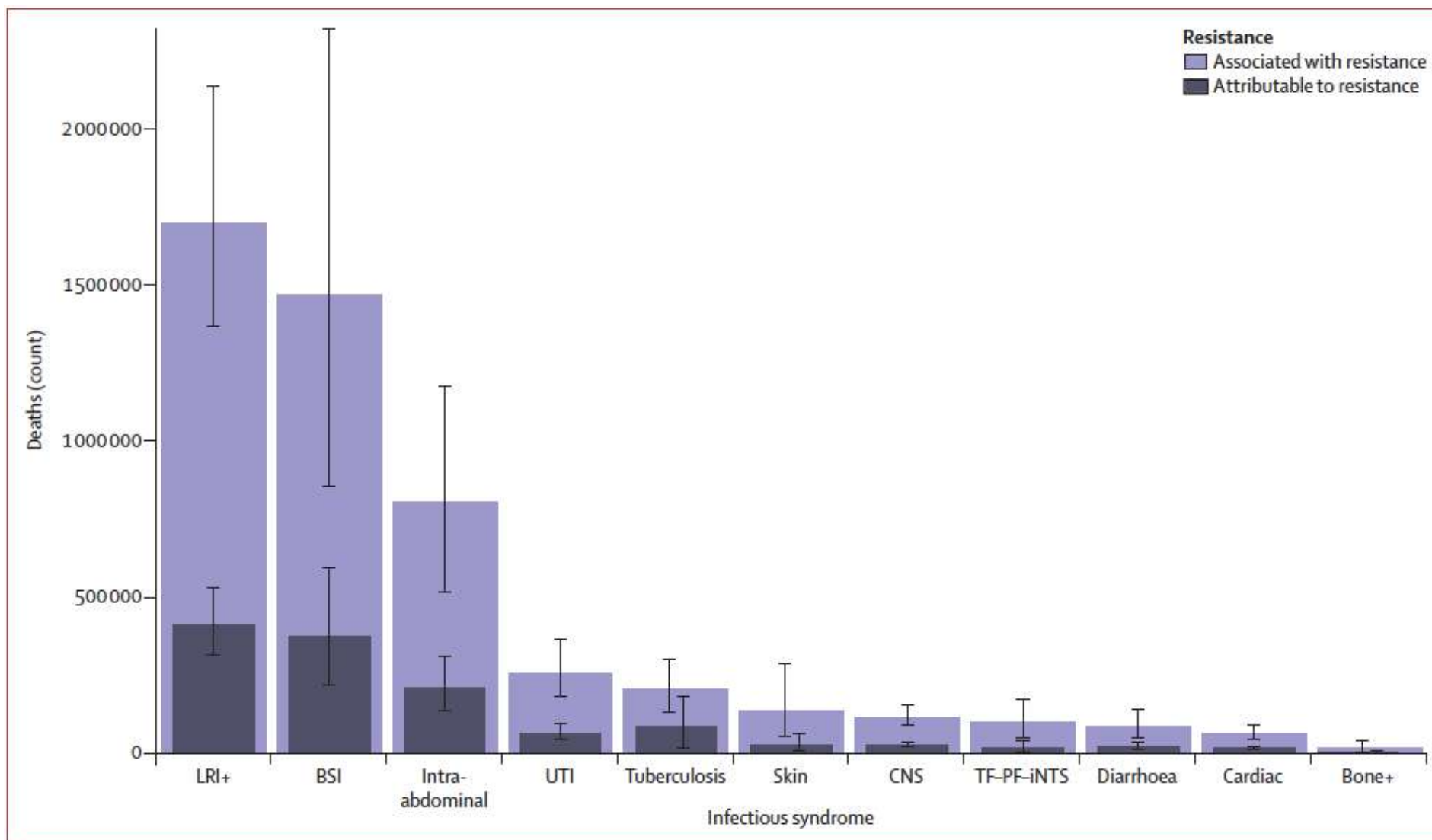


Figure 3: Global deaths (counts) attributable to and associated with bacterial antimicrobial resistance by infectious syndrome, 2019

Estimates were aggregated across drugs, accounting for the co-occurrence of resistance to multiple drugs. Error bars show 95% uncertainty intervals. Does not include gonorrhoea and chlamydia because we did not estimate the fatal burden of this infectious syndrome. Bone+=infections of bones, joints, and related organs. BSI=bloodstream infections. Cardiac=endocarditis and other cardiac infections. CNS=meningitis and other bacterial CNS infections. Intra-abdominal=peritoneal and intra-abdominal infections. LRI+=lower respiratory infections and all related infections in the thorax. Skin=bacterial infections of the skin and subcutaneous systems. TF-PF-iNTS= typhoid fever, paratyphoid fever, and invasive non-typhoidal *Salmonella* spp. UTI=urinary tract infections and pyelonephritis.

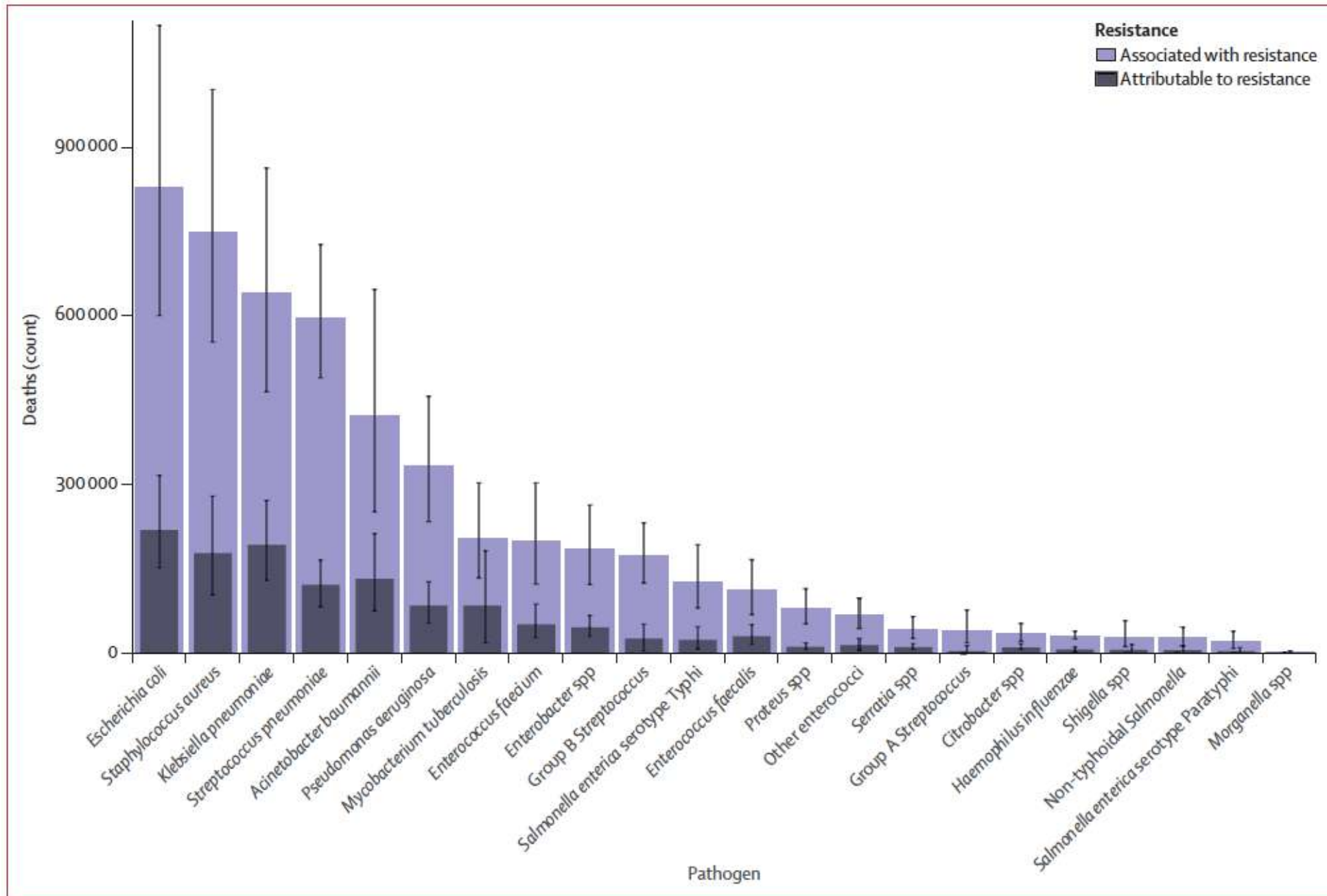


Figure 4: Global deaths (counts) attributable to and associated with bacterial antimicrobial resistance by pathogen, 2019

KOÇ UNIVERSITY Estimates were aggregated across drugs, accounting for the co-occurrence of resistance to multiple drugs. Error bars show 95% uncertainty intervals.

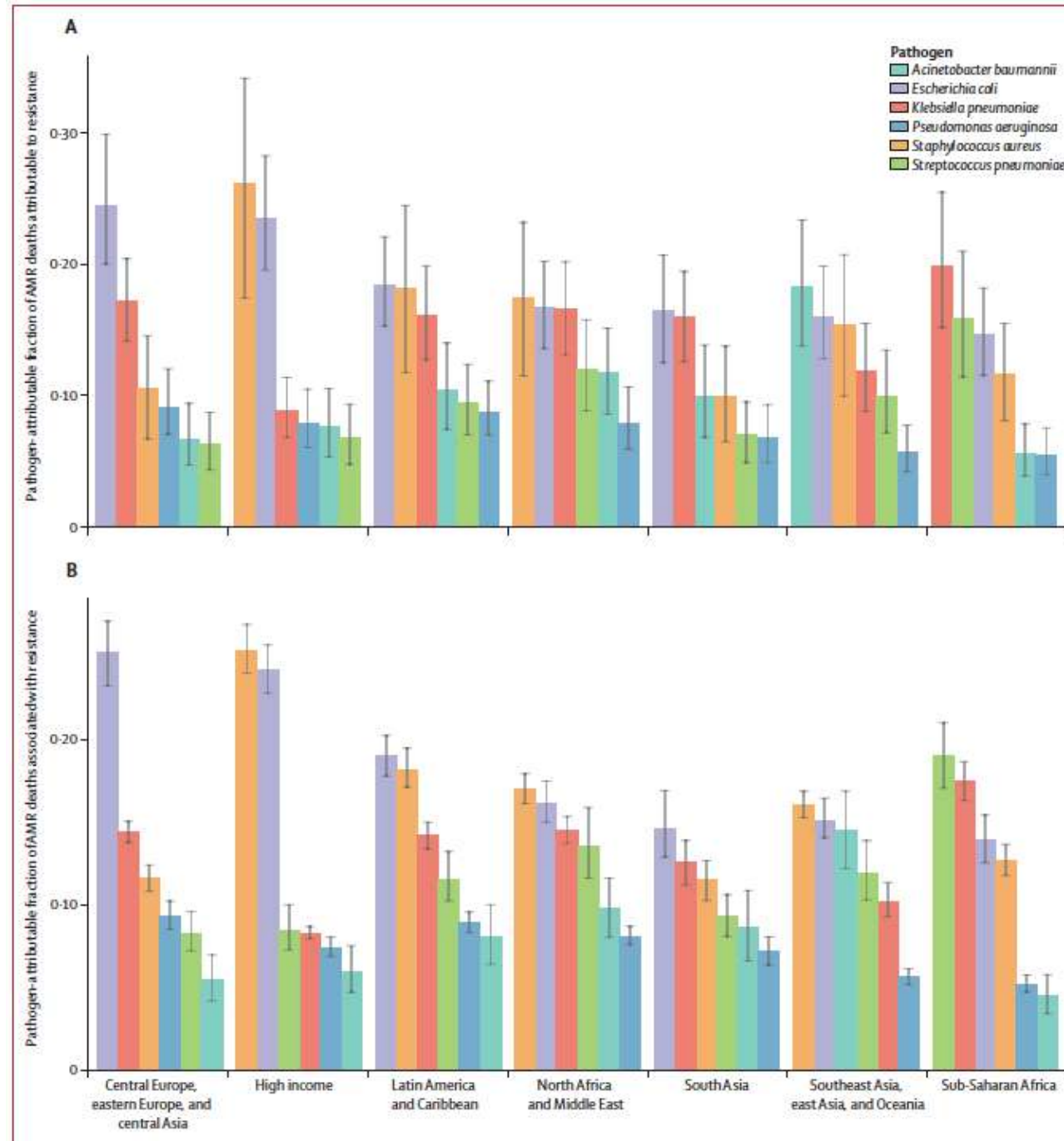
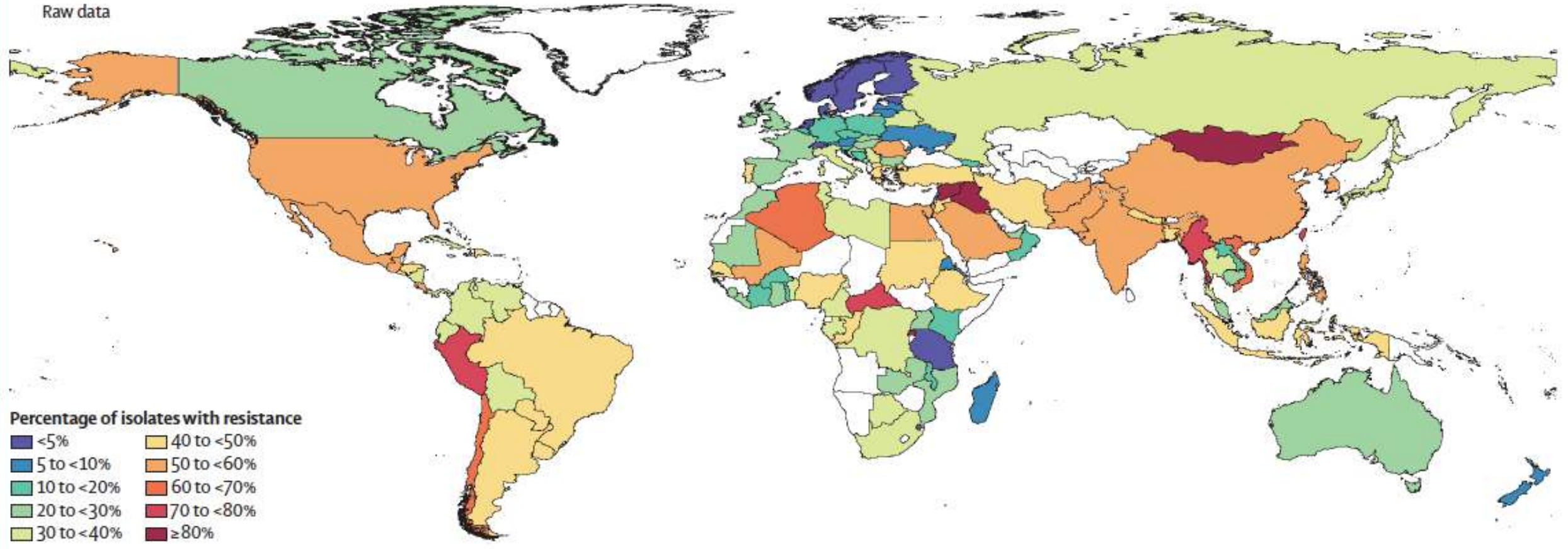


Figure 5: Pathogen-attributable fraction of deaths attributable to (A) and associated with (B) bacterial AMR for the six leading pathogens by GBD super-region, 2019



A Meticillin-resistant *Staphylococcus aureus*

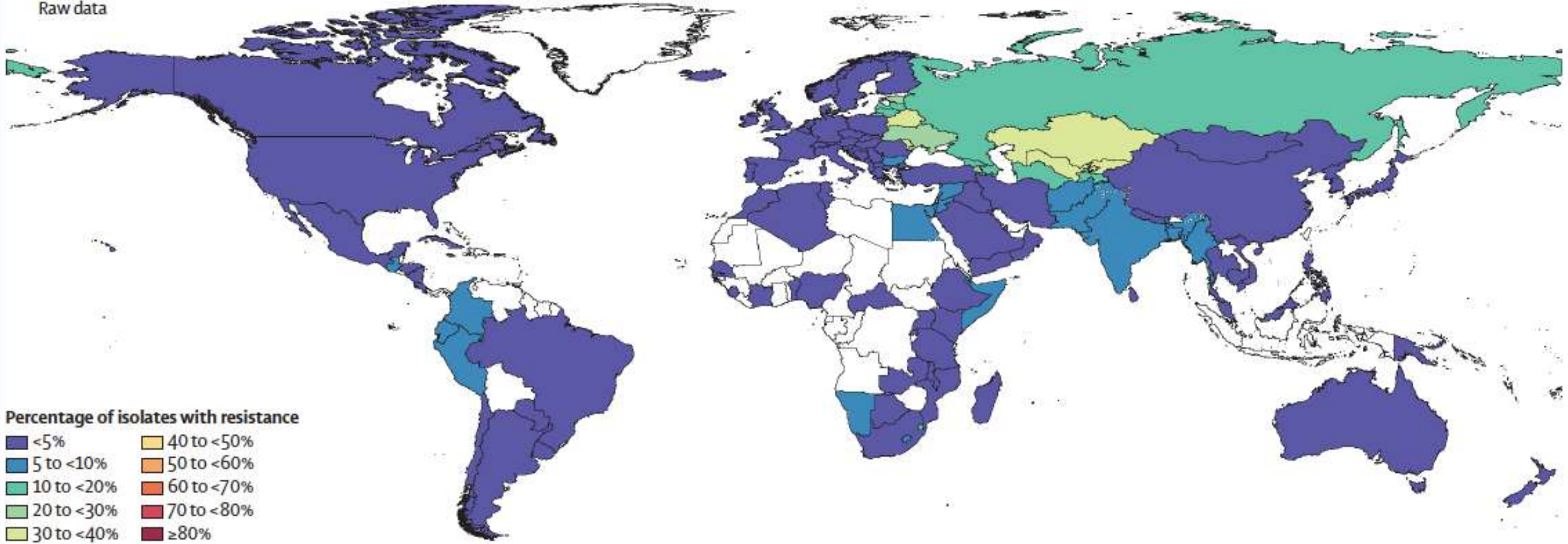
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B Isoniazid and rifampicin co-resistant (excluding XDR) *Mycobacterium tuberculosis*

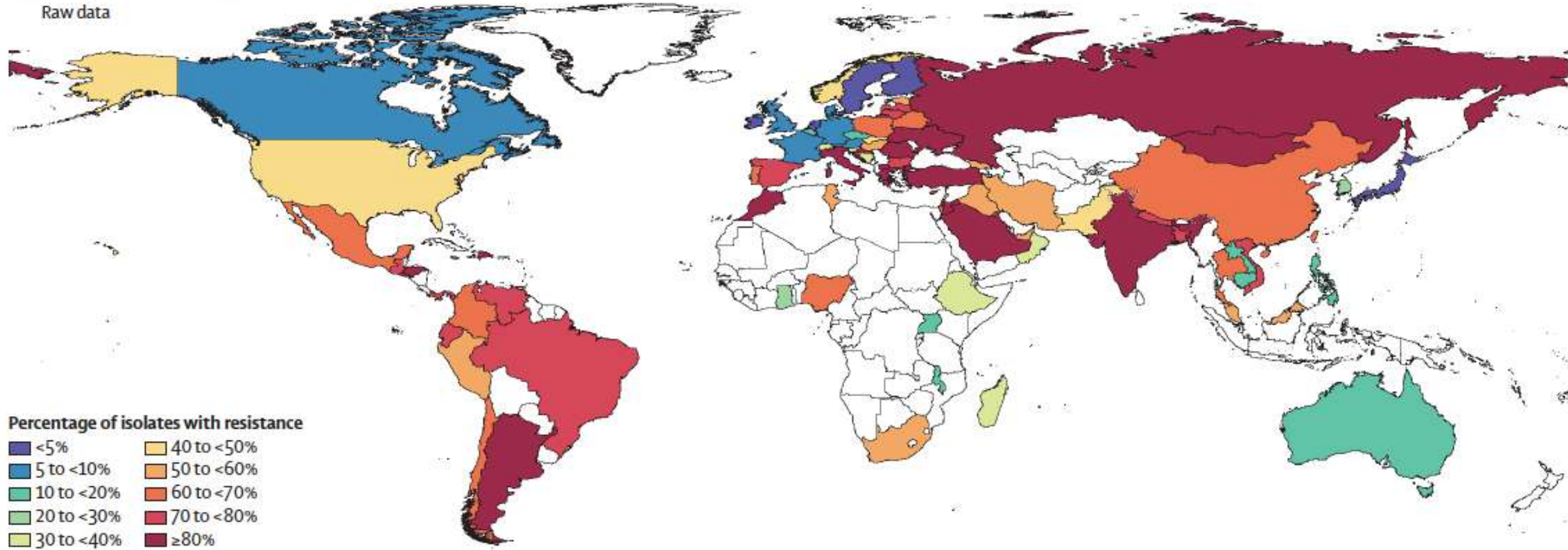
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D Carbapenem-resistant *Acinetobacter baumannii*

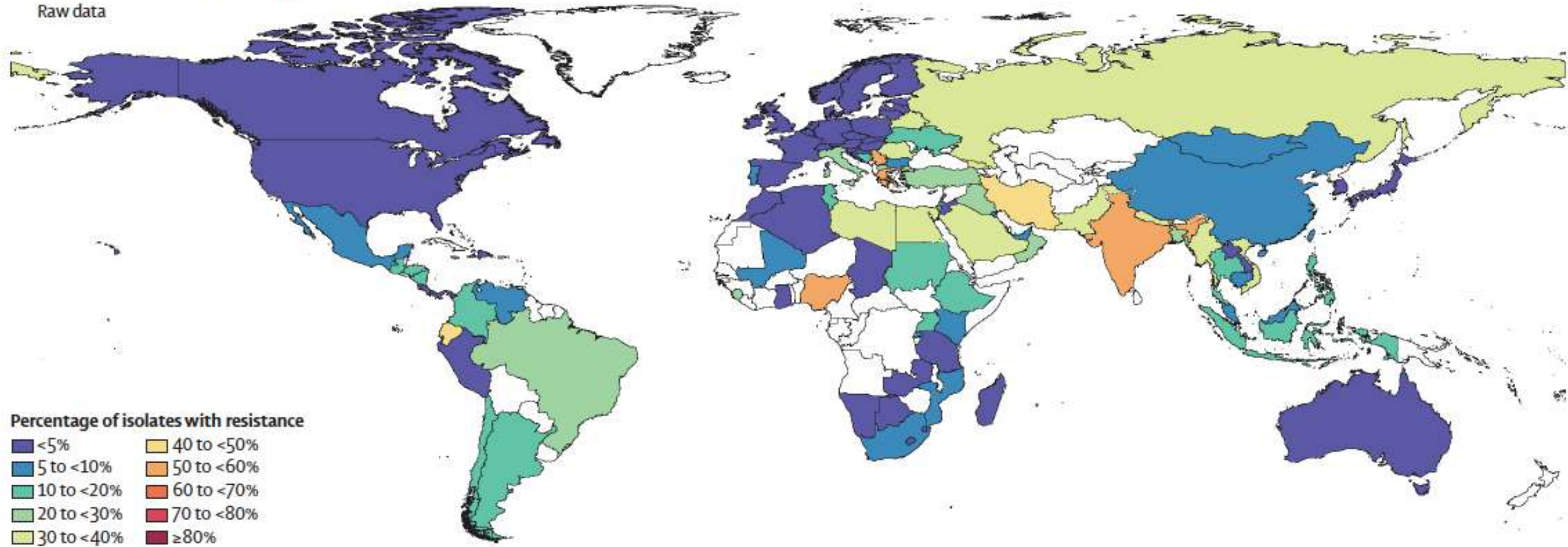
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F Carbapenem-resistant *Klebsiella pneumoniae*

Raw data





OPEN

The association between *Acinetobacter baumannii* infections and the COVID-19 pandemic in an intensive care unit

Jale Boral^{1,2,8}, Zeliha Genç^{3,8}, Fatihan Pınarlık^{1,2}, Güz Ekinci^{1,2}, Mert A. Kuskucu^{2,4}, Pelin İrkören³, Mahir Kapmaz³, Süda Tekin⁵, Nahit Çakar⁶, Evren Şentürk⁶, Fatma Yurdakul⁶, Bilge Dikenelli⁷, Fusun Can^{1,2} & Onder Ergonul^{1,2}✉



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- 1 Ocak 2018- 30 haziran 2022 arası.
- 5718 YBÜ hastası.
- 81 hastada *Acinetobacter baumannii* saptanmış.
- Pandemi döneminde önceki dönemi göre 1.9 kat artış var.
- Öncesi sonrası arasında mortalite farkı yok.

	Before <i>A. baumannii</i> outbreak	During <i>A. baumannii</i> outbreak
Training of healthcare workers		
PPE training	After recruitment	On daily basis
Preparation of cleaning solutions	After recruitment	On daily basis
Hand hygiene score	70%	97%
Glove usage	Double glove	Single glove
Cleaning procedures		
Types of cleaning solutions	Peracetic acid solution (2.0%) or chloride solution (0.1%)	Only chloride solution (0.1%)
Aspiration jars	Cleaned with surface wiping	Soaked in chloride solution
Cleaning routine	Single cleaning	Double cleaning
Ventilator related precautions		
Appropriate ventilator cleaning procedures	Standard cleaning procedures for ventilators	Separate procedures for each device
Ventilator cleaning	Ventilator cleaning	Ventilator disinfection
Transport ventilator filters	Inhalation port filter was changed	Filters for both inhalation and exhalation ports were changed
Environmental measures		
Environmental screening	None	<i>A. baumannii</i> infected rooms
Clonality surveillance	None	PFGE
Isolation of COVID-19 patients in rooms 1–8	Yes	Yes

Table 1. Infection control measures before and after *A. baumannii* outbreak.

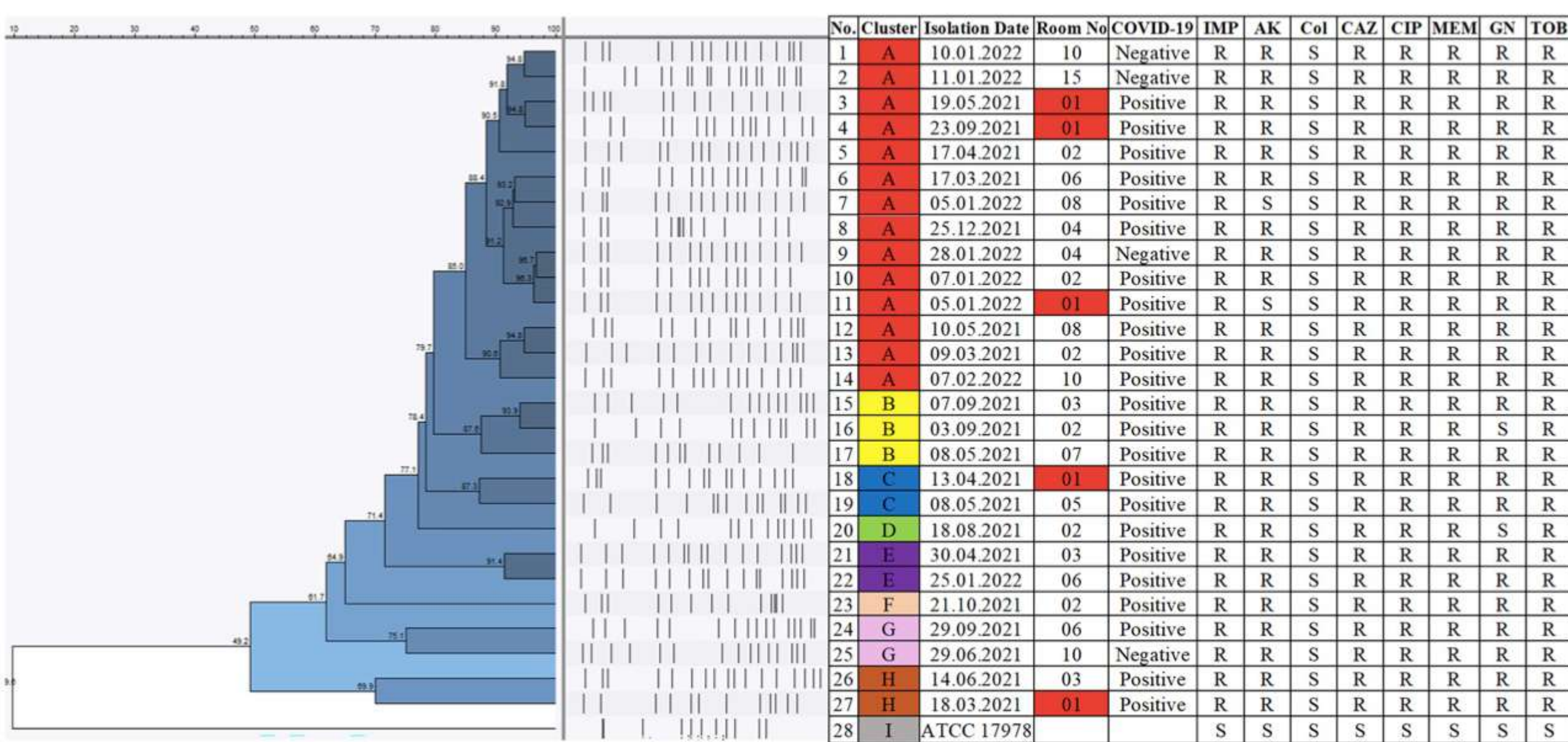


Figure 1. Cluster analysis of *A. baumannii* samples using PFGE results. Each color represents a different cluster of isolates. IMP, imipenem; AK, amikacin; Col, colistin; CAZ, ceftazidime; CIP, ciprofloxacin; MEM, meropenem; GN, gentamicin; TOB, tobramycin; R, resistant; S, susceptible.

Antimicrobial	Range	MIC50	MIC90	Susceptible (%)	Intermediate (%)	Resistant (%)
Amikacin	≤ 2 to ≥ 64	≥ 64	≥ 64	16 (19.75)	0.00	65 (80.25)
Gentamicin	≤ 1 to ≥ 16	≥ 16	≥ 16	16 (19.75)	0.00	65 (80.25)
Meropenem	≤ 0.25 to ≥ 16	≥ 16	≥ 16	8 (9.88)	0.00	73 (90.12)
Imipenem	≤ 0.25 to ≥ 16	≥ 16	≥ 16	8 (9.88)	0.00	73 (90.12)
Ceftazidime	2 to ≥ 64	≥ 64	≥ 64	7 (8.64)	0.00	74 (91.36)
Ciprofloxacin	≤ 0.25 to ≥ 4	≥ 4	≥ 4	3 (3.70)	5 (6.17)	73 (90.12)
Colistin	≤ 0.5 to 1	≤ 0.5	≤ 0.5	81 (100.00)	0.00	0.00
Piperacillin Tazobactam	≤ 4 to ≥ 128	≥ 128	≥ 128	7 (8.64)	0.00	74 (91.36)

Table 3. MIC50 and MIC90 distributions of all isolates with resistance characteristics.



Explosive COVID-19 outbreak in a German nursing home and the possible role of the air ventilation system

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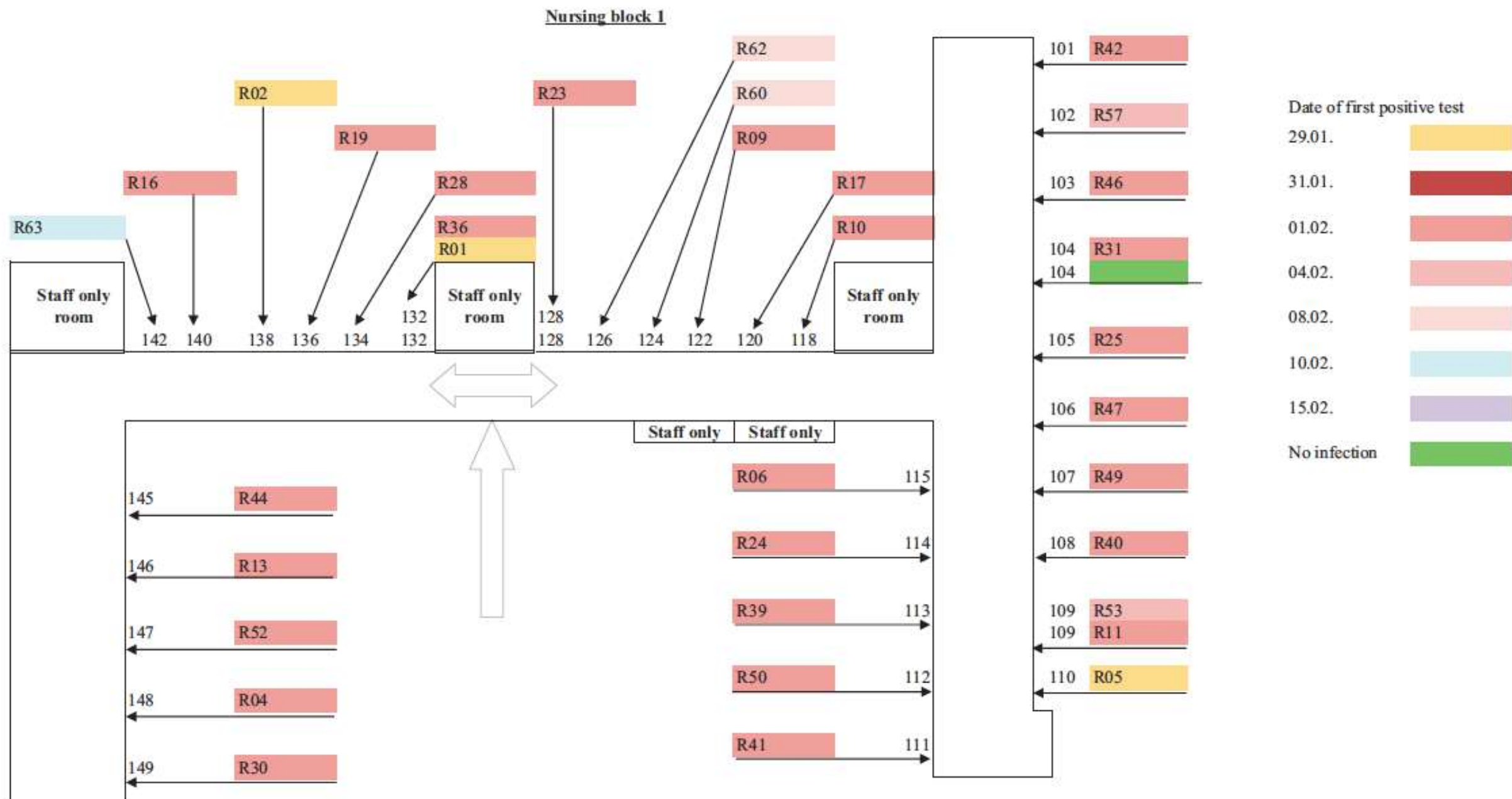
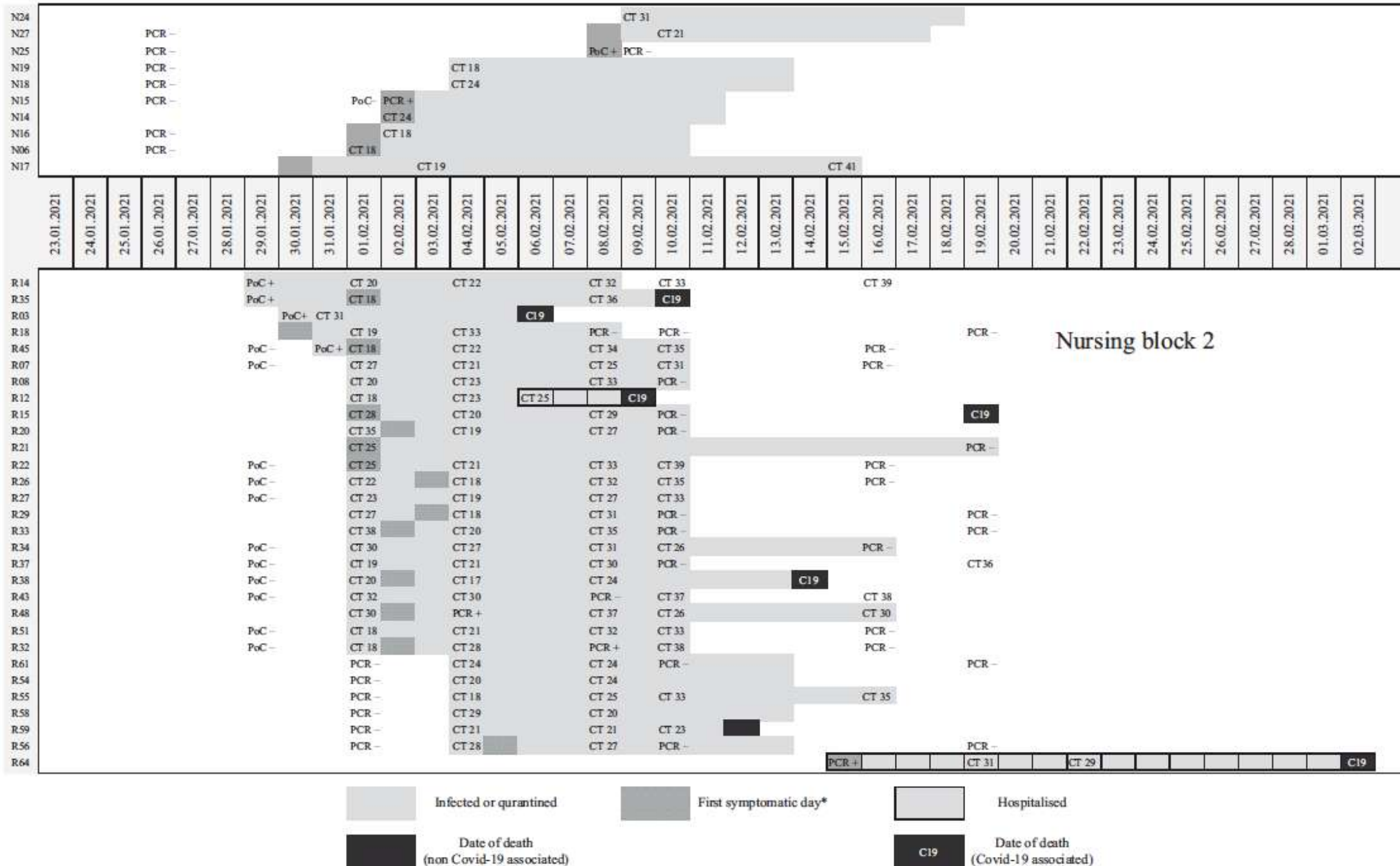


Figure 1. Floor plan of nursing block 1 with rooms 101–149; dates of COVID-19 detection are provided for each resident. R, resident.



*The first symptomatic day was not recorded for all residents



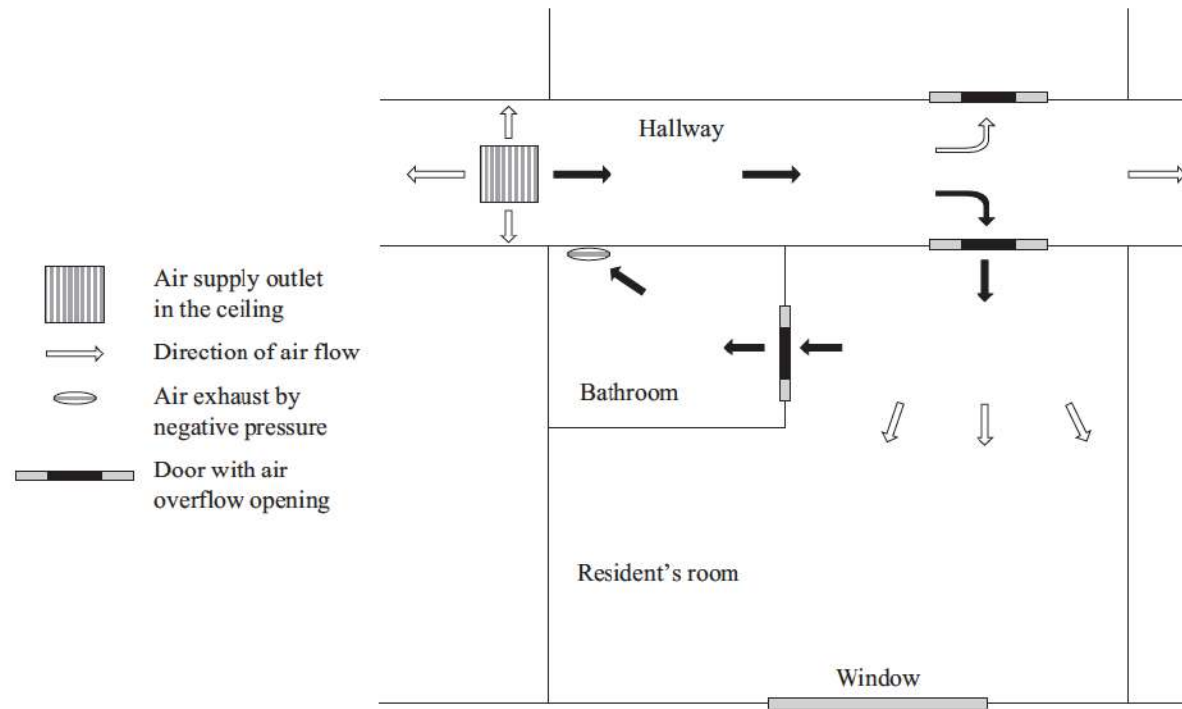


Figure 3. Schematic drawing of the air flow in the residential home; black arrows indicate the shortest possible air circulation from the air supply outlet in the ceiling of the corridor to the air exhaust in the bathroom of the resident's room (negative pressure); white arrows indicate additional directions of air flow.



Figure 4. Exhaust air opening in the tower at the highest point of the roof building of the nursing home as well as the location of the supply air opening (red arrow).

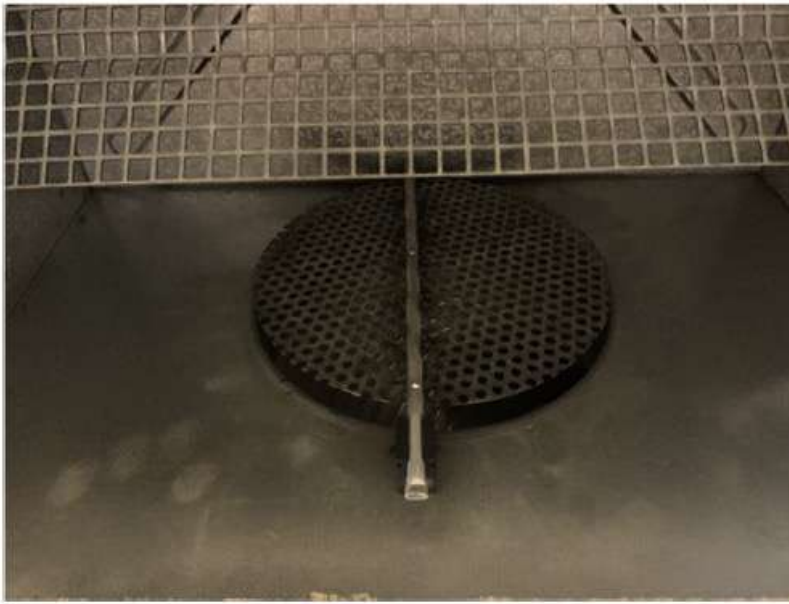


Figure 5. Grids between the air ducts and the air outlets.

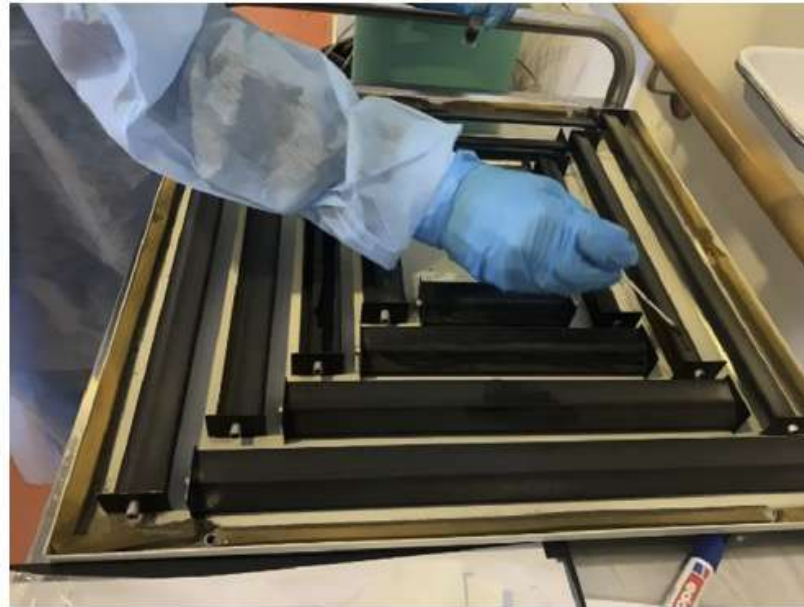


Figure 6. Sampling of the back of the air outlet.

Healthcare
Symptoms
Non-hospitalised
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COMMENTARY | ARTICLES IN PRESS

Omicron variant: assessing the duration of viral shedding and its implications

Muge Cevik, MD   • Andre C. Kalil, MD, MPH 

Published: November 24, 2022 • DOI: <https://doi.org/10.1016/j.cmi.2022.11.019>



Day 0

Viral culture

Symptoms

Symptoms

Total

References

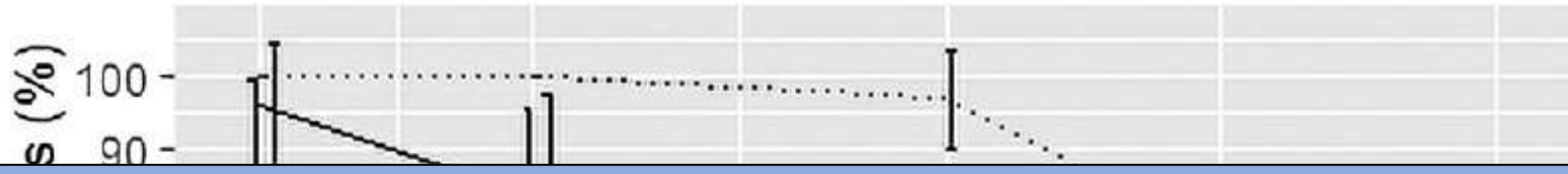
Article Info

Linked Article

Related Articles

The Omicron variant was first reported in South Africa on November 24, 2021 and has now spread to many countries [1]. Early evidence suggested reduced neutralisation antibody response of vaccines against infection with Omicron [2], raising a critical question about the infectiousness period following vaccination.

In this Clinical Microbiology and Infection edition, Keske and colleagues report the risk of infectious viral shedding with the Omicron BA.1 variant in non-hospitalised healthcare workers (HCW) in Turkey [3]. For this single-centre observational study,



- Aşılı, bağışıklığı normal olan ve Omicron ile infekte sağlık çalışanlarında 7 güne kadar bulaştırıcılık sürebilir.
- Hızlı antijen testi ve RT-PCR testi 7 güne kadar olan izolasyonun kısaltılmasında kullanılabilir.
- Düşük CT değerleri viral kültür sonucunu tahmin etmede faydalı olabilir..

---- Culture --- PCR (Ct≤24) PCR (Ct≤35) — RAT

Figure 2. Real-time reverse-transcription polymerase chain reaction, rapid antigen test, and viral culture positivity (with 95% confidence interval) by days since symptom onset in persons infected with severe acute respiratory syndrome coronavirus 2 Omicron variant. Abbreviations: Ct, cycle threshold; RAT, rapid antigen test; RT-PCR, real-time reverse-transcription polymerase chain reaction.



Incidence, Risk Factors, and Prognosis of Bloodstream Infections in COVID-19 Patients in Intensive Care: A Single-Center Observational Study

Journal of Intensive Care Medicine
2022, Vol. 37(10) 1353-1362
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Ahmet Furkan Ku
Oktay Demirkiran
Olca Dilken, MD¹
Nese Saltoğlu, MD

Table 4. Univariable and Multivariable Analyses of Risk Factors for BSIs in COVID-19

Risk factors	Unadjusted HR (95% CI)
Length of stay before ICU admission (day)	0.93 (0.90-0.97)
CRRT	2.55 (1.90-3.42)
ECMO	2.05 (0.96-4.37)
MP and TCZ	2.10 (1.04-4.24)

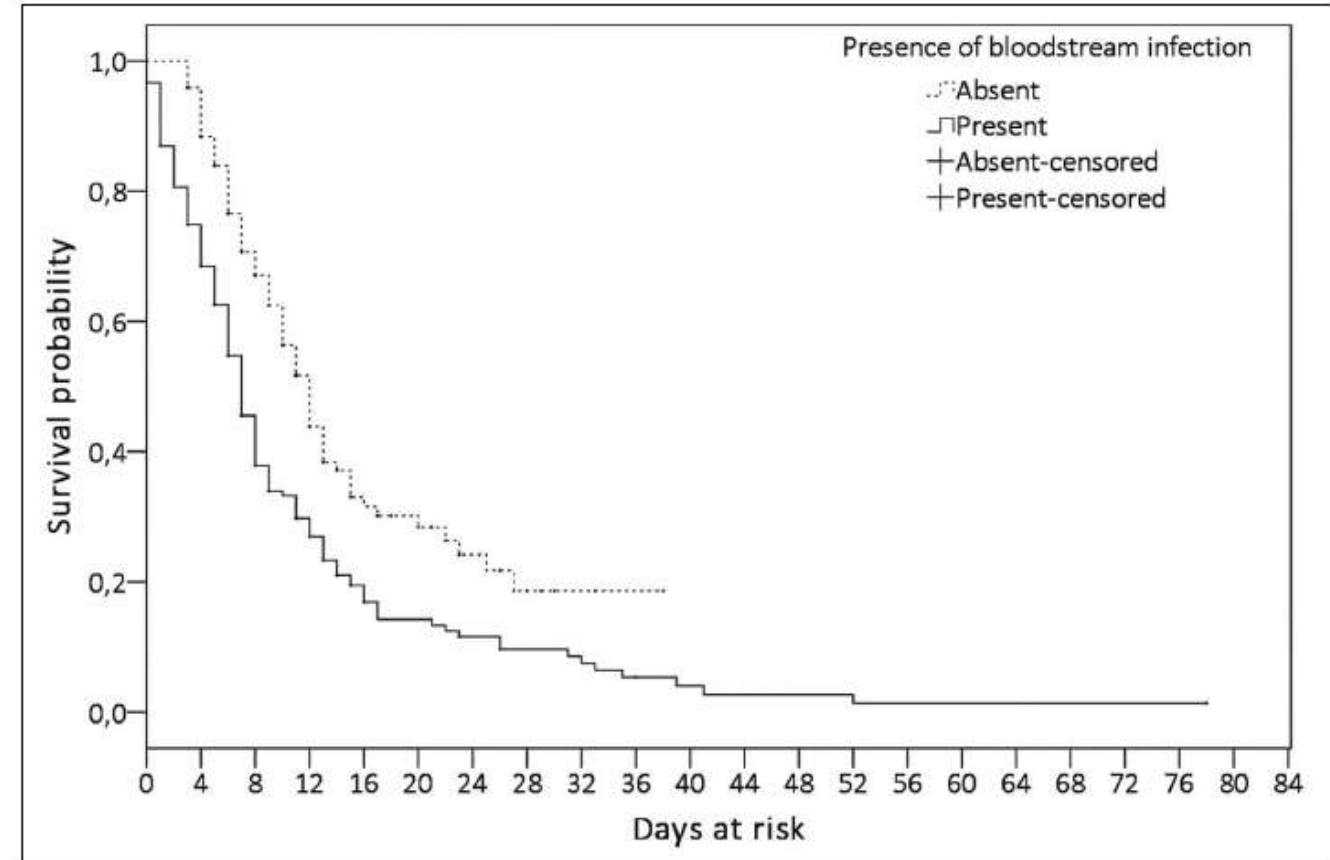


Figure 2. Comparison of the survival of the patients in terms of the presence of ICU-acquired BSI. Survival of the ICU patients with COVID-19 was analyzed via the Kaplan-Meier method. Solid line presents the survival of the patients with ICU-acquired BSI, and dotted line presents those without ICU acquired BSI. Right censoring was the discharge from the ICU. The maximum follow-up period was 38 days in the patients with ICU-acquired BSI and 78 days in the patients without ICU-acquired BSI. The origin of follow-up period started on the first day of ICU admission for the patients did not develop ICU-acquired BSI, and the first day of the first ICU-acquired BSI episode for the patients developed ICU-acquired BSI.



Major Article

A long-lasting *Sphingomonas paucimobilis* outbreak: A potential pathogens to persist on environmental devices despite disinfect measures

ay MD^c, Halide Oğuş M^e
e Çelik RN^a, Barış Otlu

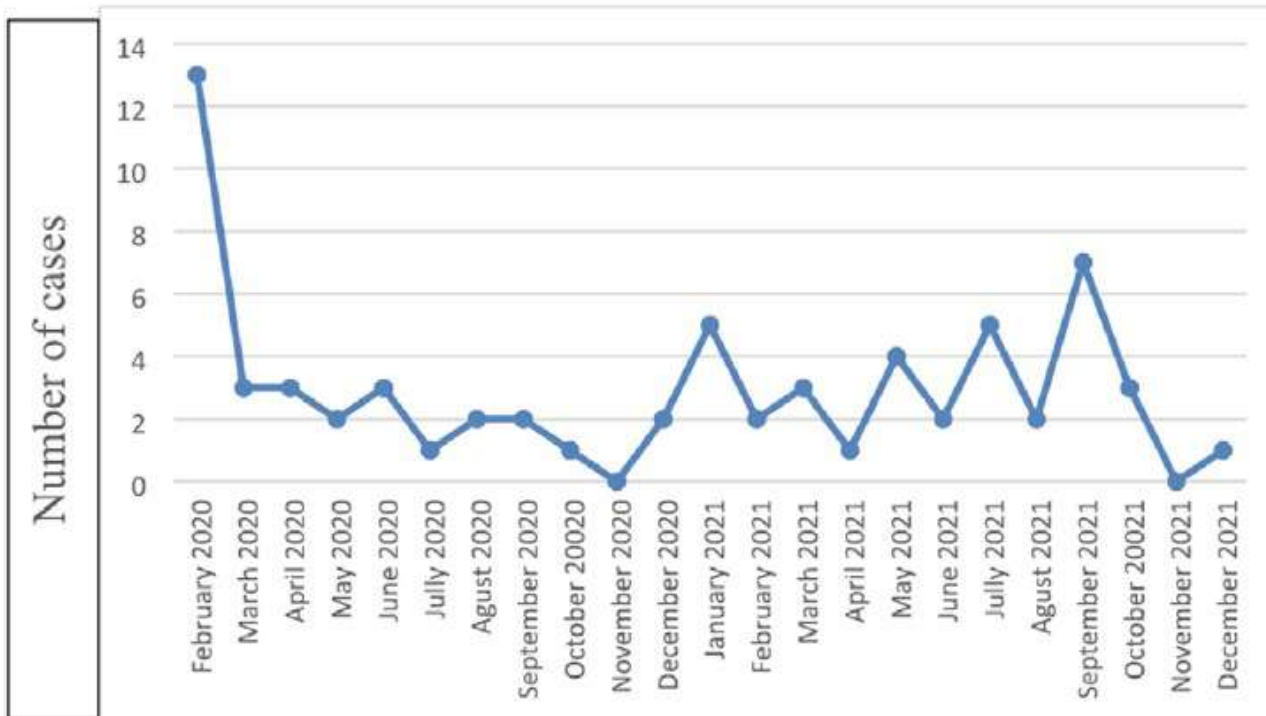
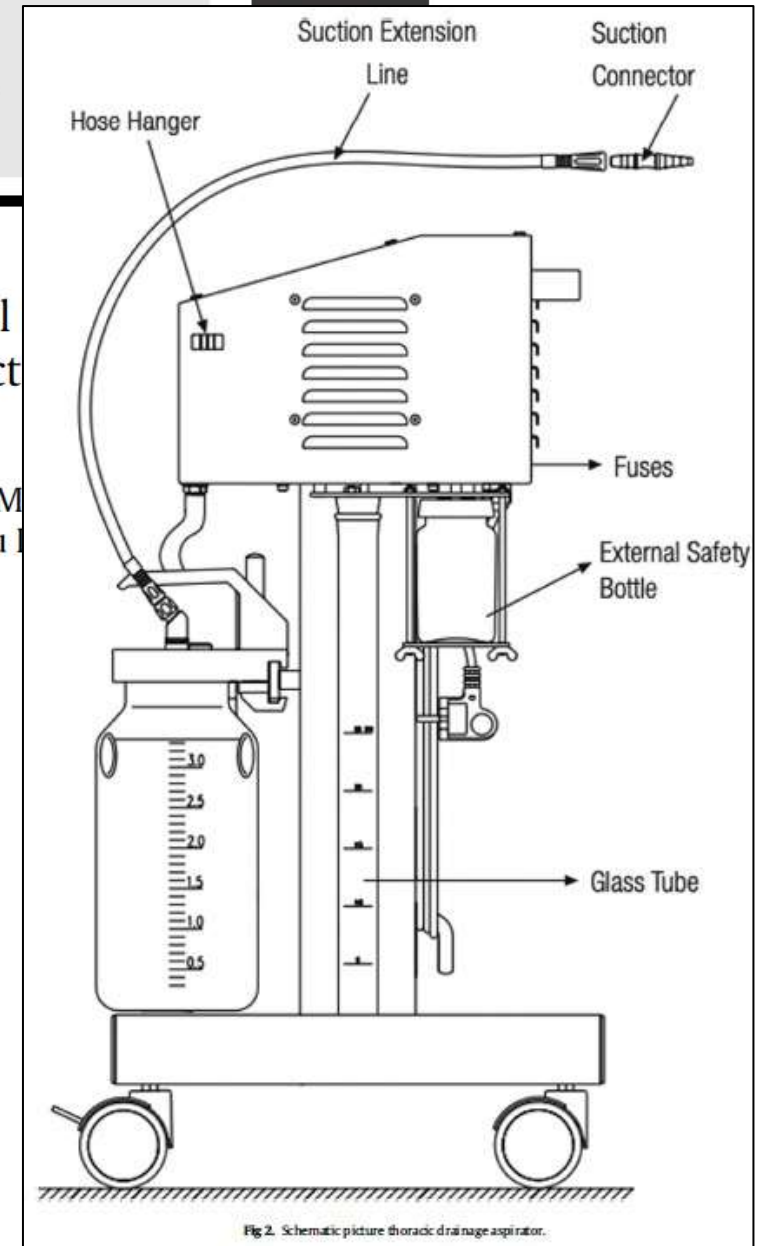
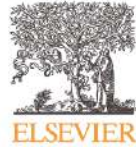


Fig 1. Distribution of *Sphingomonas paucimobilis* strains.





Major Article

Stenotrophomonas maltophilia outbreak with a commercial blood gas injector as the culprit and interventions for source and prevention: A

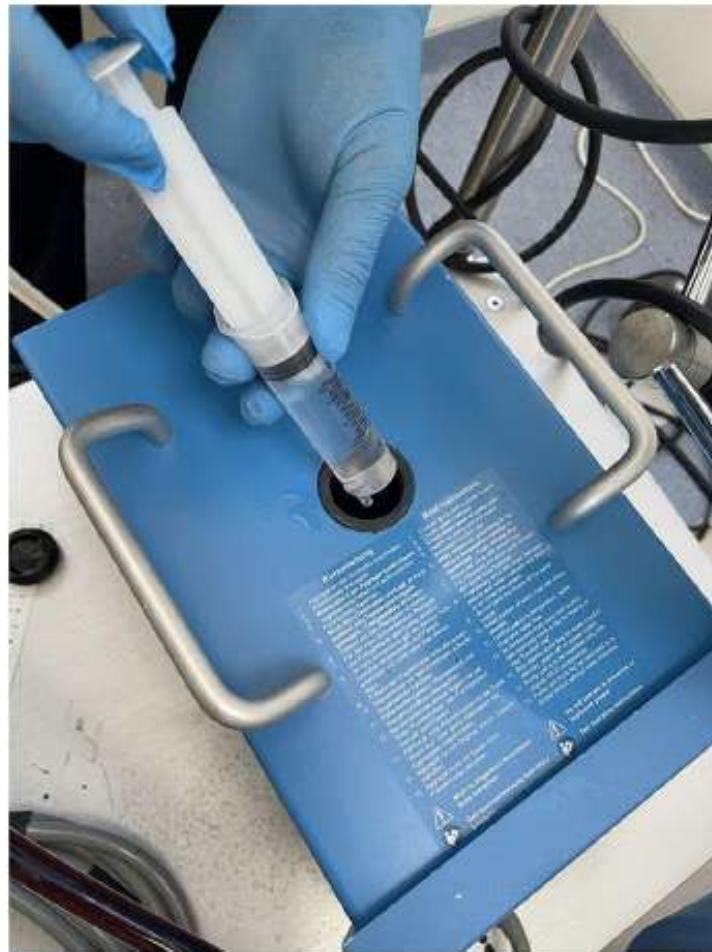


Fig 1. ECMO water heater and sampling.

W
gu
dg



Fig 2. An unused blood gas injector.



(a)

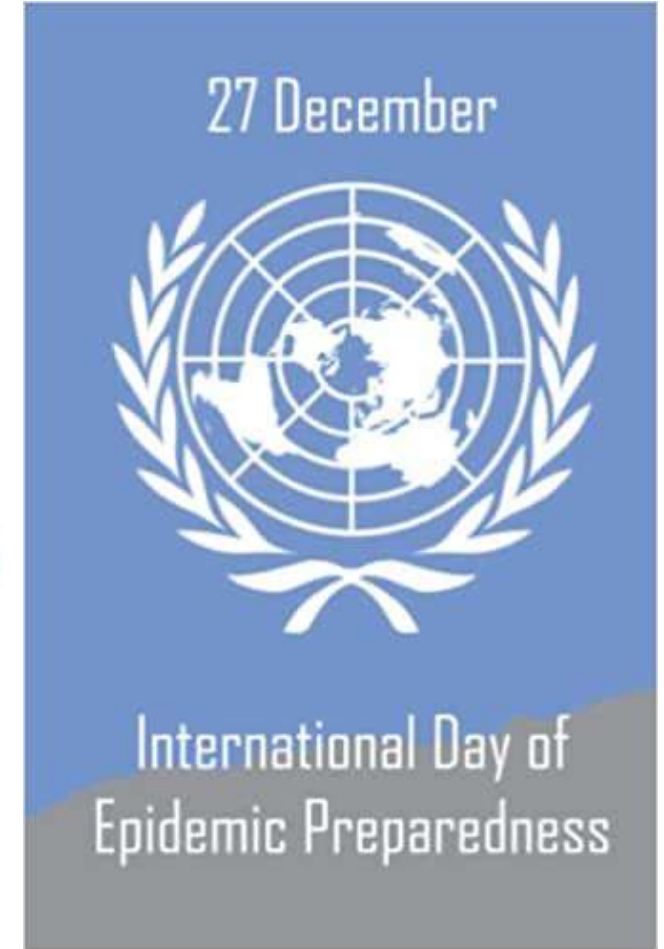


(b)

Sampling from the (a) outlet and (b) inlet lines of the ECMO oxygenator.



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International Day
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