Recent data from the European Antimicrobial Resistance Surveillance Network (EARS-Net) show a large increase of +57% in *Acinetobacter* species bloodstream infections in the European Union and European Economic Area in the first years of the COVID-19 pandemic (2020–2021) compared with 2018–2019. Most were resistant to carbapenems, from intensive care units, and in countries with ≥50% carbapenem resistance in *Acinetobacter* spp. in 2018–2019. This highlights the requirement for reinforced *Acinetobacter* preparedness and infection prevention and control in Europe.

Bloodstream infections (BSIs) with *Acinetobacter* species commonly have poor outcomes, especially in intensive care unit (ICU) patients [1]. *Acinetobacter* spp. is intrinsically resistant to many antimicrobials, and additional acquired resistance further complicates the treatment of serious infections in already vulnerable patient groups. Recent data from the European Antimicrobial Resistance Surveillance Network (EARS-Net) show a large and statistically significant increase in reports of *Acinetobacter* spp. BSIs in the European Union (EU) and European Economic Area (EEA) during the period from 2017 to 2021 [2]. Most of this increase occurred in 2020 and 2021, the first years of the coronavirus disease (COVID-19) pandemic. Here we further explore this trend in a subset of data from laboratories that continuously reported data during that period.

**Data description**

Our data originate from qualitative routine antimicrobial susceptibility testing (AST) results of blood isolates collected by local clinical laboratories in national networks in EU/EEA countries. These results are reported annually by national centres to the European Centre for Disease Prevention and Control (ECDC), according to the EARS-Net reporting protocol [3]. In its analyses, EARS-Net only includes the first isolate per patient each year and for each bacterial species.

All EU countries, Iceland and Norway reported data to EARS-Net every year during the period 2017 to 2021 [2,4]. For this analysis, we restricted the dataset to BSIs with *Acinetobacter* spp. and to only those laboratories that reported carbapenem (imipenem and/or meropenem) antimicrobial susceptibility testing results for *Acinetobacter* spp. for every year in 2017 to 2021 (255 of 826 laboratories reporting, on average, per year). We made this restriction to limit bias from year-to-year changes in the number, hospital affiliation and type of reporting laboratories, and because not all countries can discriminate between laboratories that did not report and those that had no cases. The United Kingdom ceased reporting data to ECDC in 2020 when it withdrew from the EU and was hence not included. In addition, France was excluded because, following a major reorganisation of national surveillance in 2020, only a few laboratories were continuously identifiable. The Table presents data for the 28 included countries, with and without restriction to continuously reporting laboratories.

As the resistance percentages for *Acinetobacter* spp. varied substantially between EU/EEA countries [4], we grouped the countries according to their mean national annual carbapenem resistance percentage in
### Table

Annual carbapenem-resistant *Acinetobacter* species bloodstream infections and carbapenem susceptibility testing results for *Acinetobacter* species bloodstream infections, in all laboratories that reported data to EARS-Net and in those that continuously reported data, EU/EEA countries, 2017–2021 (n = 31,242 infections)

| Carbapenem susceptibility test result | 2017 | 2018 | 2019 | 2020 | 2021 | Number of cases in 2020–2021 vs 2018–2019 | Number of cases in 2020 vs average 2017–2019 | Number of cases in 2021 vs average 2017–2019 |
|--------------------------------------|------|------|------|------|------|------------------------------------------|---------------------------------------------|---------------------------------------------|
|                                      | n    | %    | n    | %    | n    | % Change | p value<sup>c</sup> | % Change | p value<sup>c</sup> | % Change | p value<sup>c</sup> |
| All laboratories (annual mean = 826 laboratories)<sup>a</sup> | | | | | | | | | | | |
| R                                   | 2,831 | 59.2 | 3,105 | 60.1 | 2,629 | 56.5 | 4,379 | 65.2 | 7,396 | 74.5 | +105.4 | <0.001<sup>d</sup> | +53.4 | <0.001<sup>d</sup> | +159.1 | <0.001<sup>d</sup> |
| S/I                                 | 1,950 | 40.8 | 2,063 | 39.9 | 2,028 | 43.5 | 2,333 | 34.8 | 2,528 | 25.5 | +18.8 | <0.001<sup>d</sup> | +15.9 | <0.001<sup>d</sup> | +25.5 | <0.001<sup>d</sup> |
| All                                 | 4,781 | 100.0 | 5,168 | 100.0 | 4,657 | 100.0 | 6,712 | 100.0 | 9,924 | 100.0 | +69.3 | <0.001<sup>d</sup> | +37.9 | <0.001<sup>d</sup> | +103.8 | <0.001<sup>d</sup> |
| Continuously reporting laboratories (n = 255 laboratories)<sup>b</sup> | | | | | | | | | | | |
| R                                   | 1,237 | 48.5 | 1,293 | 48.3 | 1,354 | 48.4 | 1,891 | 57.6 | 3,767 | 70.8 | +113.8 | <0.001<sup>d</sup> | +46.1 | <0.001<sup>d</sup> | +191.0 | <0.001<sup>d</sup> |
| R (Group 1 countries<sup>e</sup>) | 30    | 3.6  | 37    | 4.4  | 17    | 1.9  | 29    | 3.3  | 23    | 2.3  | −3.7  | 0.85  | +3.6  | 0.90  | −17.9 | 0.49 |
| R (Group 2 countries<sup>e</sup>) | 60    | 35.3 | 38    | 21.5 | 26    | 20.6 | 30    | 22.7 | 104   | 49.8 | +109.4 | <0.001<sup>d</sup> | −27.4 | 0.18 | +151.6 | <0.001<sup>d</sup> |
| R (Group 3 countries<sup>e</sup>) | 1,147 | 74.6 | 1,218 | 73.5 | 1,311 | 72.9 | 1,832 | 80.6 | 3,640 | 88.3 | +116.4 | <0.001<sup>d</sup> | +49.5 | <0.001<sup>d</sup> | +197.1 | <0.001<sup>d</sup> |
| S/I                                 | 1,314 | 51.5 | 1,382 | 51.7 | 1,444 | 51.6 | 1,390 | 42.4 | 1,554 | 29.2 | +4.2  | 0.12  | +0.7  | 0.85  | +12.6 | 0.001 |
| S/I/R                               | 2,551 | 100.0 | 2,675 | 100.0 | 2,798 | 100.0 | 3,281 | 100.0 | 5,321 | 100.0 | +57.2 | <0.001<sup>d</sup> | +22.7 | <0.001<sup>d</sup> | +98.9 | <0.001<sup>d</sup> |

EARS-Net: European Antimicrobial Resistance Surveillance Network; EEA: European Economic Area; EU: European Union; I: susceptible, increased exposure; R: resistant; S: susceptible, standard dosing regimen.

<sup>a</sup> Data for France and the United Kingdom were excluded.

<sup>b</sup> Only includes EU/EEA laboratories that were identifiable as having reported ≥ 1 Acinetobacter spp. isolate with carbapenem susceptibility data every year in 2017–2021.

<sup>c</sup> Poisson regression model to assess the statistical significance of changes in the numbers of bloodstream infections.

<sup>d</sup> The chi-squared test comparing the percentage of carbapenem resistance in 2020–2021 vs 2018–2019 also had p < 0.001.

<sup>e</sup> The reporting countries were grouped according to the mean of their crude, national, annual percentage of Acinetobacter spp. resistance to carbapenems in 2018 and 2019. These were Group 1 (< 10% carbapenem resistance in 2018–2019): Austria, Belgium, Denmark, Estonia, Finland, Germany, Iceland, Ireland, Luxembourg, Malta, the Netherlands, Norway and Sweden; Group 2 (10% to < 50% carbapenem resistance in 2018–2019): Czechia, Portugal, and Slovenia; Group 3 (≥ 50% carbapenem resistance in 2018–2019): Bulgaria, Croatia, Cyprus, Greece, Hungary, Italy, Latvia, Lithuania, Poland, Romania, Slovakia and Spain. Percentages in these rows refer to carbapenem-resistant isolates among all isolates analysed for the countries in a given group. The raw data for the group level are provided in the Supplement.
Countries in Group 1 had < 10% carbapenem resistance (n = 13; Austria, Belgium, Denmark, Estonia, Finland, Germany, Iceland, Ireland, Luxembourg, Malta, the Netherlands, Norway, Sweden), Group 2 had 10% to < 50% carbapenem resistance (n = 3: Czechia, Portugal, Slovenia) and Group 3 had ≥ 50% carbapenem resistance (n = 12: Bulgaria, Croatia, Cyprus, Greece, Hungary, Italy, Latvia, Lithuania, Poland, Romania, Slovakia, Spain). When stratifying by patient ward type, we grouped the units as ‘ICU’ (adult and paediatric ICUs), ‘not ICU’ (all other ward types) and ‘unknown ward type’ (no information available on ward type).

We assessed the statistical significance of changes in the numbers of BSIs and in the percentage of carbapenem resistance comparing 2020–2021 with 2018–2019, using Stata Statistical Software (Release 15.1. College Station, TX: StataCorp LLC) for a Poisson regression model and a chi-squared test, respectively, with p values < 0.05 considered as significant.

The observed trends for Acinetobacter spp. BSI in the EU/EEA are worrying because resistance to carbapenems causes a high burden of disease in vulnerable hospitalised patients [5-7]. Our findings suggest that countries where carbapenem-resistant Acinetobacter spp. were already well established before the COVID-19 pandemic (Group 3) had the biggest challenges in controlling further spread in 2020–2021.

Acinetobacter spp. is difficult to eradicate from the hospital environment, colonising hospital patients and staff and causing outbreaks, particularly in ICUs [1]. Several reports have identified Acinetobacter spp. as one of the most frequent causes of infectious complications in hospitalised patients with COVID-19 [8-10]. The observed increasing trends at EU/EEA level compared with the pre-pandemic situation [2,11,12] were probably driven by the profound impact of the COVID-19 pandemic on hospital care, which increased the number of patients at risk of Acinetobacter spp. BSI, and also by difficulties in applying infection prevention and control (IPC) measures. In 2020–2021, there were larger numbers of severely ill patients, many with severe pulmonary infection. High occupancy rates necessitated increased provision of ICU beds, often with staff who were overworked or less experienced [13,14]. Inappropriate application of
Contact precautions for COVID-19 patients, in particular suboptimal hand hygiene, as well as contamination and insufficient cleaning of the hospital environment, probably contributed to direct or indirect between-patient *Acinetobacter* spp. transmission [15-20]. Finally, reduced attention to antimicrobial stewardship, with resulting increased carbapenem use, may have contributed [21].

For context, in 2020–2021 compared with 2018–2019, the laboratories that continuously reported *Acinetobacter* spp. data to EARS-Net also reported more cases of BSI with *Enterococcus faecium* (+29%), *E. faecalis* (+16%), *Pseudomonas aeruginosa* (+8%), *Klebsiella pneumoniae* (+6%), but these differences were much less pronounced than for BSI with *Acinetobacter* spp. (+57%). Laboratories reported fewer cases of BSI with *Streptococcus pneumoniae* (−47%), *Escherichia coli* (−5%) and *Staphylococcus aureus* (−1%). These differences probably depend on the epidemiological characteristics of the various pathogens. For example, *S. pneumoniae* and *E. coli* are more frequently transmitted in the community and in non-ICU hospital settings. During the COVID-19 pandemic, transmission of microorganisms in the community was affected by containment actions such as stay-at-home orders, physical distancing, hygiene measures and the use of face masks. This may have contributed to the sharp decline in typically community-acquired infections such as those caused by *S. pneumoniae* [22,23].

There were exceptions to the general trends by country group, indicating that the trends were not only explain by the pre-pandemic percentage of carbapenem resistance. For example, Portugal and Spain were outliers in their respective groups by reporting fewer *Acinetobacter* spp. BSIs than in 2018–2019, whereas Slovenia reported a larger increase in *Acinetobacter* spp. BSIs than other Group 2 countries.

Although reasons for the trends observed during the COVID-19 pandemic remain to be clarified, most factors that potentially favoured the increase in carbapenem-resistant *Acinetobacter* spp. infections, and in general multidrug-resistant microorganisms, are amenable to public health intervention. Options include rigorous adherence to hand hygiene, environmental cleaning, provision and appropriate

**Figure 2**

Bloodstream infections with carbapenem-resistant *Acinetobacter* species, reported by laboratories that continuously reported data to EARS-Net, by country group* and year, EU/EEA, 2017–2021 (n = 9,542)

- **Group 1** (13 countries): Austria, Belgium, Denmark, Estonia, Finland, Germany, Iceland, Ireland, Luxembourg, Malta, the Netherlands, Norway and Sweden.
- **Group 2** (3 countries): Czechia, Portugal, and Slovenia.
- **Group 3** (12 countries): Bulgaria, Croatia, Cyprus, Greece, Hungary, Italy, Latvia, Lithuania, Poland, Romania, Slovakia and Spain.

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*The reporting countries were grouped according to the mean of their crude, national, annual percentage of *Acinetobacter* spp. resistance to carbapenems in 2018 and 2019. These were Group 1 (<10% carbapenem resistance in 2018–2019): Austria, Belgium, Denmark, Estonia, Finland, Germany, Iceland, Ireland, Luxembourg, Malta, the Netherlands, Norway and Sweden; Group 2 (10% to <50% carbapenem resistance in 2018–2019): Czechia, Portugal, and Slovenia; Group 3 (≥50% carbapenem resistance in 2018–2019): Bulgaria, Croatia, Cyprus, Greece, Hungary, Italy, Latvia, Lithuania, Poland, Romania, Slovakia and Spain.*

EARS-Net: European Antimicrobial Resistance Surveillance Network; EEA: European Economic Area; EU: European Union.
The large increase in carbapenem-resistant *Acinetobacter* spp. BSI in the EU/EEA during a time of great challenges for healthcare calls for reinforced application of the preparedness and response actions that we present above. Surveillance at local, national and EU/EEA levels will be vital to monitor whether this worrying development is halted or even reversed.

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**Figure 3**

Percentage and number of bloodstream infections with carbapenem-resistant *Acinetobacter* species from laboratories that continuously reported data to EARS-Net, by country group*, EU/EEA, 2018–2019 vs 2020–2021 (n = 9,542)

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EARS-Net: European Antimicrobial Resistance Surveillance Network; EEA: European Economic Area; EU: European Union.

* The reporting countries were grouped according to the mean of their crude, national, annual percentage of *Acinetobacter* spp. resistance to carbapenems in 2018 and 2019. These were Group 1 (<10% carbapenem resistance in 2018–2019): Austria, Belgium, Denmark, Estonia, Finland, Germany, Iceland, Ireland, Luxembourg, Malta, the Netherlands, Norway and Sweden; Group 2 (10% to <50% carbapenem resistance in 2018–2019): Czechia, Portugal, and Slovenia; Group 3 (≥50% carbapenem resistance in 2018–2019): Bulgaria, Croatia, Cyprus, Greece, Hungary, Italy, Latvia, Lithuania, Poland, Romania, Slovakia and Spain.

* Reported <25 *Acinetobacter* spp. isolates in both periods 2018–2019 and 2020–2021. As the data are restricted to laboratories that reported continuously in 2017–2021, the percentages and numbers for countries may differ from the dataset that included all laboratories (Table) [2,4].

For Latvia, the two datapoints for the mean overlap and only the dark symbol is visible.
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This manuscript presents a subset of data in an online database 'the ECDC surveillance atlas of infectious diseases' (ECDC Atlas) [4]. Although the ECDC Atlas contains the full dataset, users cannot generate the restricted dataset presented in our manuscript. Similarly, an upcoming ECDC publication, intended for publication on 17 November 2022 for European Antibiotic Awareness Day (EAAD; 18 November 2022), contains the full dataset, and a different restricted dataset. It is the 'Antimicrobial resistance in the EU/EEA (EARS-Net) - Annual Epidemiological Report for 2021' [2].

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Conflict of interest

None declared.

Authors’ contributions

Conceptualisation and design of the study, and writing of the first draft: CG, DLM, HM, LDH, PK. Initial interpretation of the results and revision of the draft: CG, DLM, DP, HM, LDH, PK. The EARS-Net Study Group, CG, DLM, DP, HM, LDH and PK contributed to acquisition and analysis of the data and interpretation of results, subsequently critically reviewing the final manuscript.

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Ethical statement

This study only included anonymised surveillance data; therefore, ethical approval was not required.

Data availability statement

This manuscript presents a subset of data in an online database 'the ECDC surveillance atlas of infectious diseases' (ECDC Atlas) [4]. Although the ECDC Atlas contains the full dataset, users cannot generate the restricted dataset presented in our manuscript. Similarly, an upcoming ECDC publication, intended for publication on 17 November 2022 for European Antibiotic Awareness Day (EAAD; 18 November 2022), contains the full dataset, and a different restricted dataset. It is the 'Antimicrobial resistance in the EU/EEA (EARS-Net) - Annual Epidemiological Report for 2021' [2].
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