

# Understanding the impact of resistant bugs on deaths in England

<b>Submission date</b> 28/05/2024	<b>Recruitment status</b> Recruiting	<input checked="" type="checkbox"/> Prospectively registered <input checked="" type="checkbox"/> Protocol
<b>Registration date</b> 02/06/2024	<b>Overall study status</b> Ongoing	<input checked="" type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
<b>Last Edited</b> 14/10/2025	<b>Condition category</b> Other	<input type="checkbox"/> Individual participant data <input checked="" type="checkbox"/> Record updated in last year

## Plain English summary of protocol

### Background and study aims

Antimicrobial resistance refers to when bacteria, viruses, fungi and parasites (bugs) no longer respond to antibiotic medicines. It is usually caused by prolonged and repeated exposure of bugs to antibiotics, in people, animals or the environment, which helps bugs eventually build defence mechanisms against them. These bugs can then spread via contact from person to person, causing resistant infections to people who have not had antibiotics before. For this reason, previously effective treatments stop working due to antimicrobial resistance worldwide. Research has shown that infections from antibiotic resistant bugs are more likely to lead to death, disability, and additional stay in hospital. It has also been estimated that, by 2050, 10 million deaths a year will be caused by antimicrobial resistance.

Despite the importance of antimicrobial resistance, which has been characterised as a global health emergency by the World Health Organisation, we have limited information about how many people die every year due to infections resistant to antibiotics. This is because of a variety of factors. For example, studies have shown that doctors are more likely to record chronic health problems like cancer in the death certificate as the primary cause of death, and the contribution of infection is often underestimated. Another problem is that in order to find antibiotic resistant bugs, cultures from the patient need to be taken, which is not always performed. This leads to underestimation of true numbers of antibiotic resistant infections. Yet, knowing exactly how many people die due to antimicrobial resistance is important, as it drives political and public awareness about the problem, highlighting the need for better treatments.

Very few previous studies have tried to estimate the true numbers of deaths due to antibiotic resistant infections. They have primarily used a technique called modelling to do so. Modelling uses math to estimate the numbers of deaths, based on how groups of patients with antibiotic sensitive and antibiotic resistant infections behave, as well as total numbers of infections. Because modelling doesn't look at individual patient cases though, it can lead to erroneous results and underestimation of the problem.

In this study, we will attempt for the first time to calculate the total number of deaths caused by antibiotic resistant infections in England in 2021, 2022 and 2023. We aim to do that by anonymously linking the cause of death of each patient as documented on the death certificate

with the bugs they were positive for in cultures and specimens up to 28 days before the date of death.

Who can participate?

Every patient who died in England in 2021, 2022 and 2023 will be included in the study but their data will be collected in a way that will not allow them to be identified. Participants who expressed their wish for their data not to be used for research purposes before they died will be excluded. We believe that with this approach we will get more accurate data on the total number of people in England, in whom antimicrobial resistance contributed to their death.

What does the study involve?

Retrospective analysis of patient records.

What are the possible benefits and risks of participating?

None

Where is the study run from?

University College London (UK)

When is the study starting and how long is it expected to run for?

March 2024 to June 2026

Who is funding the study?

UK Health Security Agency

Who is the main contact?

ioannis.baltas.20@ucl.ac.uk

l.grandjean@ucl.ac.uk

## Contact information

### Type(s)

Principal investigator

### Contact name

Dr Louis Grandjean

### ORCID ID

<https://orcid.org/0000-0002-1457-8327>

### Contact details

UCL GOS Institute of Child Health, Infection, Immunity & Inflammation Department, 30 Guilford St

London

United Kingdom

WC1N 1EH

+44 7712336582

l.grandjean@ucl.ac.uk

### Type(s)

Public, Scientific, Principal investigator

**Contact name**

Dr Ioannis Baltas

**ORCID ID**

<https://orcid.org/0000-0002-8008-4835>

**Contact details**

UCL GOS Institute of Child Health, Infection, Immunity & Inflammation Department, 30 Guilford St

London

United Kingdom

WC1N 1EH

+44 7543780069

ioannis.baltas.20@ucl.ac.uk

## **Additional identifiers**

**Clinical Trials Information System (CTIS)**

Nil Known

**Integrated Research Application System (IRAS)**

340243

**ClinicalTrials.gov (NCT)**

Nil known

**Protocol serial number**

IRAS 340243, CPMS 61354

## **Study information**

**Scientific Title**

Recording Antimicrobial Resistance during Death Certification in England

**Acronym**

AMR-DC

**Study objectives**

1. That it is possible to calculate AMR-associated deaths by linking death certification and patient-level microbiological data.
2. That AMR-associated mortality has been increasing in England during the study period.
3. That AMR was associated with a significant number of deaths (>2% of total deaths) in England in 2021-2023.
4. That a significant proportion of deaths (>10%) due to infection in England in 2021, 2022 and 2023 were AMR-associated.
5. That ESBLs were the AMR resistance mechanism with the highest number of AMR-associated deaths in England during the study period.
6. That most AMR-associated deaths (>90%) occur in hospital, disproportionately affect patients in Intensive Care and patients under Haematology and Oncology.

7. That patients recording AMR-associated deaths have longer hospital length of stays, are more likely to be admitted to Intensive Care and require more prolonged organ support.

### **Ethics approval required**

Ethics approval required

### **Ethics approval(s)**

approved 20/03/2024, North West - Haydock Research Ethics Committee (2 Redman Place, Stratford, London, E20 1JQ, United Kingdom; +44 (0)207 104 8032; haydock.rec@hra.nhs.uk), ref: 24/NW/0084

### **Study design**

Retrospective observational cohort

### **Primary study design**

Observational

### **Study type(s)**

Other

### **Health condition(s) or problem(s) studied**

Antimicrobial resistance effect on mortality

### **Interventions**

This will be a retrospective observational cohort study. Eligible patients will be identified through the Civil Registrations of Death database provided by NHS England and will be linked to the HES database, also held by NHS England, to obtain additional clinical metadata. Pseudonymised cases will be subsequently linked to the SGSS database (AMR module) held by UKHSA in order to detect clinical samples with AMR pathogens of interest within 28 days of the patient's death. Anonymised results will be analysed using a standardized pathway to determine whether each death was AMR-associated. The rate and total number of deaths associated with AMR in this cohort, the most important pathogens, pathogen drug-combinations and resistance mechanisms will be described.

### **Intervention Type**

Other

### **Primary outcome(s)**

Measured using clinical databases:

The total number of patients with AMR-associated deaths in England in 2021-2023

### **Key secondary outcome(s)**

Measured using clinical databases:

1. The percentage of AMR-associated deaths in 2021, 2022 and 2023, including the monthly trend of totals numbers.
2. The total number and percentage of AMR-associated deaths for each site of infection, AMR pathogen of interest, pathogen-drug combination, and resistance mechanism.
3. Underlying cause of death (UCODs) and Multiple cause of death (MCDs) groupings percentages for the entire study cohort.
4. The total number and percentage of UCOD and MCDs that were AMR-associated.
5. The total number and percentage of patients with AMR-associated deaths who die in hospital

versus in the community.

6. The percentage of AMR-associated deaths per medical specialty for patients who died in hospital.

7. The association between AMR and healthcare utilisation (mode of admission, duration of stay in hospital, ICU admission rates and duration of stay, life-support duration).

8. The total number and percentage of patients with AMR pathogens of interest in sterile sites within 28 days of death without infection being recorded in the pathway that led to death in the MCCD.

9. The total number of patients who had AMR documented in their death certificate (by using the ICD-10 codes U82-84).

### **Completion date**

30/06/2026

## **Eligibility**

### **Key inclusion criteria**

All patients who had their death registered in England between 01/01/2021 and 31/12/2023 will be included in this study.

### **Participant type(s)**

Patient

### **Healthy volunteers allowed**

No

### **Age group**

All

### **Lower age limit**

0 years

### **Upper age limit**

150 years

### **Sex**

All

### **Total final enrolment**

1520000

### **Key exclusion criteria**

Only patients who had expressed the wish for their data not to be used for research purposes under the National Data Opt-Out (DOO) will be excluded.

### **Date of first enrolment**

01/07/2024

### **Date of final enrolment**

31/03/2026

# Locations

## Countries of recruitment

United Kingdom

England

## Study participating centre

**UCL Great Ormond Street Institute of Child Health**

30 Guilford Street

London

United Kingdom

WC1N 1EH

# Sponsor information

## Organisation

University College London

## ROR

<https://ror.org/02jx3x895>

# Funder(s)

## Funder type

Government

## Funder Name

UK Health Security Agency

# Results and Publications

## Individual participant data (IPD) sharing plan

The datasets generated during and/or analysed during the current study will be stored for 3 years and may be available upon request from Dr Louis Grandjean ([l.grandjean@ucl.ac.uk](mailto:l.grandjean@ucl.ac.uk))

## IPD sharing plan summary

Available on request

## Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
<a href="#">Participant information sheet</a>	Participant information sheet	11/11/2025	11/11/2025	No	Yes
<a href="#">Protocol file</a>	version 1.1	25/05/2024	29/05/2024	Yes	No
<a href="#">Protocol file</a>	version 1.2	08/10/2024	30/10/2024	No	No
<a href="#">Statistical Analysis Plan</a>	Section 5.8 of the protocol version 1.1	25/05/2024	10/06/2024	No	No
<a href="#">Study website</a>	Study website	11/11/2025	11/11/2025	No	Yes