# Using hospital network-based surveillance for antimicrobial resistance as a more robust alternative to self-reporting

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19

## 20 Abstract

Hospital performance is often measured using self-reported statistics, such as the incidence of hospital-transmitted micro-organisms or those exhibiting antimicrobial resistance (AMR),

23 encouraging hospitals with high levels to improve their performance. However, hospitals that

24 increase screening efforts will appear to have a higher incidence and perform poorly, undermining

25 comparison between hospitals and disincentivising testing, thus hampering infection control. We

26 propose a surveillance system in which hospitals test patients previously discharged from other

27 hospitals and report observed cases. Using NHS Hospital Episode Statistics data, we analysed

patient movements across England and assessed the number of hospitals required to participate in

such a reporting scheme to deliver robust estimates of incidence. With over 1.2 million admissions

to English hospitals previously discharged from other hospitals annually, even when only a fraction

of hospitals (41/155) participate (each screening at least 1000 of these admissions), the proposed
 surveillance system can estimate incidence across all hospitals. By reporting on other hospitals, the

reporting of incidence is separated from the task of improving own performance. Therefore the

incentives for increasing performance can be aligned to increase (rather than decrease) screening

35 efforts, thus delivering both more comparable figures on the AMR problems across hospitals and

36 improving infection control efforts.

37

38 Keywords: Antimicrobial Resistance; Surveillance; Patient sharing; Healthcare network

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#### 39 Introduction

- 40
- 41 Many healthcare systems worldwide mandate the reporting of key hospital statistics to measure
- 42 performance[1]. Such self-reported assessments are intended to provide a clear, comparable
- 43 overview of each hospital's status, by ranking them based on their reported statistics. Poorly
- 44 performing hospitals can then be encouraged to improve using incentives ranging from financial
- 45 penalties[2,3] to reputational damage through 'naming and shaming'. The mandatory reporting of
- 46 antimicrobial resistance (AMR) and other hospital-transmitted organisms are examples of
- 47 commonly used self-reporting systems[4].
- 48
- 49 Surveillance systems for AMR are attractive to policy-makers, as they can be used to increase
- 50 patient safety by identifying where extra infection prevention and control (IPC) efforts need to be
- 51 coordinated, as well as providing insight into the spread and epidemiology of AMR. Changes in
- 52 incidence after introducing such systems, like the dramatic decline in methicillin-resistant
- 53 Staphylococcus aureus (MRSA) bacteraemia after the initiation of the mandatory surveillance
- 54 scheme in the United Kingdom[5], have led some to conclude that such self-reporting surveillance
- 55 systems help reduce rates.
- 56
- 57 However, true incidence of AMR is often hard to measure, because large numbers of affected
- 58 patients may be asymptomatically colonised[6,7] and thus only found when actively screened.
- 59 Hospitals targeting screening strategies to identify more cases may thus worsen their ranking by
- 60 increasing their reported incidence. Systems of assessing hospitals based on self-reported carriage
- 61 rates may thus unintentionally punish hospitals with stringent testing, screening, and reporting
- 62 regimes, because of their seemingly poor performance. Both IPC efforts and hospital performance
- 63 monitoring may therefore be hindered by the conflicting incentives: to improve IPC efforts, a
- 64 hospital needs to identify as many cases as possible, while it needs to find as few as possible to
- 65 improve its performance ranking.
- 66
- 67 We explore how to align incentives for hospitals, by separating the task of reporting incidence of a
- 68 predominantly carried micro-organism that is acquired in hospital from the task of lowering its
- 69 incidence. To do this, we propose a novel surveillance system based on the hospital network formed
- 70 by shared patients, namely testing patients that were previously admitted to another hospital to
- 71 provide an approximation of the incidence of AMR in that hospital. We show the potential of this
- 72 network-based surveillance system to provide incidence estimates, and explore its operational
- 73 limits, in particular the number of participating hospitals needed to reliably estimate incidences for
- all hospitals. We argue that such a system can provide a more robust surveillance system for AMR
- 75 than self-reporting.
- 76

#### 77 Methods

- 78
- 79 Network-based surveillance system
- 80 In the proposed surveillance system (Figure 1), each hospital reports the number of patients
- 81 previously admitted to and discharged from other hospitals in a predefined time-frame (e.g. the
- 82 previous 12 months) and found to be colonised when screened on admission to this index hospital
- 83 (denoted imported cases). The reported numbers of imported cases are then pooled to give the total
- 84 number of found cases exported from all the hospitals across the network. For simplicity of
- 85 reporting, any untested patients are assumed to not be colonised (providing an incentive to test
- 86 admissions previously discharged from elsewhere). The number of imported cases are then divided
- 87 by the total number of patients previously discharged from that hospital and admitted to one of the
- 88 reporting hospital (which can be obtained from central statistics) to give an estimate of incidence.

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- 89 Alternatively, without loss of generality, numbers tested and testing positive could be reported and
- 90 summed to give an estimate of incidence.
- 91
- 92 To demonstrate, we use data on patient admissions from the National Health Service (NHS)
- 93 Hospital Episode Statistics (HES) to determine the number of patients that were admitted to
- 94 different English hospital Trusts (denoted 'hospitals') post discharge. We sorted all admissions per
- 95 patient by admission date; and for all admissions during 2014-'15 determined whether the previous
- 96 discharge happened 1 year before the admission date and whether the previous admission was to a
- 97 different hospital to the current one (i.e. the patient was shared between hospitals). Sensitivity
- 98 analyses considered 6, 3, 1 month and one week.
- 99
- 100 Each hospital had two sets of admissions associated with it: 1) all admissions (the general patient
- 101 population), and 2) a subset the admissions of patients previously discharged from another hospital,
- 102 now admitted to this hospital (the received patients). The received patient population comes from a
- 103 number (potentially all) of the other hospitals. We therefore denote the number of patients
- 104 discharged from hospital *i* and subsequently admitted to hospital *j* as  $m_{ij}$ , where  $s_i = \sum_j m_{ij}$  is the total
- 105 shared population size from hospital *i*. Under the proposed surveillance scheme, these received
- 106 patients should be screened as they are admitted to hospital *j* to gather information about the
- 107 incidence of hospital-associated pathogens in hospital *i*.
- 108
- 109 Coverage
- 110 The system consists of the "reporting set", namely hospitals reporting the number of AMR cases
- among their received patients, and the "covered set", namely hospitals whose discharged patients
- are screened as they arrive in other hospitals. We consider a hospital to be part of the covered set
- 113 once a fixed number of its discharged patients per year (the reporting threshold) are received by the
- 114 hospitals within the reporting set. Thus the reporting set does not necessarily need to include all
- 115 hospitals for the covered set to include all hospitals.
- 116
- 117 Any hospital sharing fewer patients than this reporting threshold with all other hospitals combined
- 118 cannot, by definition, be reported on by such a scheme. Thus the minimum number of patients
- shared by hospitals is the highest reporting threshold that can be used (n=1216). Taking 1000
- 120 shared patients as the reporting threshold, we determined the total number of hospitals that need to
- be included in the surveillance scheme to be able to report on all hospitals in three ways; first by
- 122 random assignment, second by adding hospitals based on the number of received patients, and third
- 123 by adding hospitals using a greedy algorithm.
- 124

#### 125 Assignment of hospitals

- 126 For the first selection procedure, we randomly added hospitals to the reporting set, one at a time,
- 127 calculating the number of hospitals in the covered set after each addition. Hospitals were added to
- 128 the reporting set until all hospitals were included in the covered set, repeating this algorithm 100
- 129 times. For the second procedure (receipt-based), we sorted hospitals based on the total number of
- 130 patients they received from other hospitals, and added them to the reporting set, starting with the
- 131 hospital that received most patients and iteratively adding the other hospitals to maximise the
- 132 number of received patients added at each step.
- 133
- 134 The greedy algorithm iteratively added the hospital to the reporting set that would add the most
- 135 hospitals to the covered set. Per step, we calculated for each reporting hospital how many other
- 136 hospitals it would add information on (i.e. by how many hospitals the covered set would increase if
- 137 this hospital was added to the reporting set). If the number of covered hospitals did not increase by
- 138 adding any hospital, the hospital that resulted in the largest increase in number of received patients

- 139 from hospitals not yet included in the covered set was added. The same procedure was used if two
- 140 hospitals resulted in the same increase to the covered set.
- 141
- 142 Reciprocal reporting (snow-ball effect)
- 143 We further tested the effect of assuming that covered hospitals will automatically start reporting
- 144 once they are themselves reported on, based on the game-theoretical considerations that hospitals
- 145 will try to 'win' the ranking of reported incidences (supplementary text). After adding a hospital
- 146 following the greedy algorithm, we checked if all covered hospitals were present in the reporting set
- 147 and added them if they were not. Because the increase in reporting could increase the number of
- 148 covered hospitals, this step was repeated until no hospitals were added to the reporting and covered
- 149 sets. After this, the next hospital was added to the reporting set using the greedy algorithm again.
- 150

#### 151 **Results**

- 152 Network-based surveillance
- 153 To test the feasibility of having hospitals report the number of patients previously admitted to other
- 154 hospitals that are AMR (or other equivalent carried micro-organism) positive on admission, rather
- than self-reporting their own patients colonised on or during admission, we reconstructed the
- 156 English hospital network (Figure 2A), based on the NHS Hospital Episode Statistics for England.
- 157 The network consisted of 155 hospital organisations (so-called Trusts, denoted 'hospitals' for
- 158 generalisability) during the financial year 2014-15, admitting 8,681,397 patients for a total of
- 159 15,708,764 admissions. A total of 1,208,999 admissions were preceded within a year by a discharge
- 160 from a different hospital, mainly concentrated within a small number of strong connections between
- 161 hospitals (Figure 2B). The median time between the previous discharge and admission was 28 days
- 162 (IQR 6-104), the mean number of overnight stays was 2.1 (IQR 0-2, median 0) for all patient
- admissions (Figure 2C), while shared patients stayed 4.6 nights (IQR 0-4, median 1).
- 164
- 165 The number of shared patients (patients who were first admitted to a certain hospital, and
- 166 subsequently admitted to any of the others) was highest for a tertiary care hospital in the North-East
- 167 (23,260 received by others), and lowest for a cancer centre in the North-West (1,216 received by
- 168 others). Based on 1,216 as the upper limit of patients that can be received from the least connected
- 169 hospital, we set our reporting threshold at 1000. If the maximum time between discharge and
- 170 subsequent admission was reduced from a year to a week, the number of subsequent admissions
- 171 was reduced by about 78% (Figure 3A), with a total of 264,920 subsequent admissions, of which
- 172 5,314 were received from the most-connected (a London teaching hospital) and 232 from the least-
- 173 connected (an orthopaedic hospital). Specialist hospitals shared the fewest patients, and higher
- thresholds up to 2,989 can be used to include the remaining 146 hospitals when these nine
- 175 specialists are excluded.
- 176
- 177 A key feature of this system is that hospitals can be included in the covered set even if none of the
- 178 individual reporting hospitals receive over the threshold of 1000 patients, as long as all hospitals
- 179 combined receive over this threshold. In fact, a median 134 hospitals (of total 155) were required in
- 180 a randomly chosen reporting set to provide enough data to include information about all hospitals in
- 181 the covered set. Strikingly, a median of only 30 hospitals needed to be included in a randomly
- 182 chosen reporting set to survey incidence in half (n=78) of the hospitals. Numerous hospitals
- 183 received enough patients to be able to individually report on several others (Figure 3B). Four
- 184 hospitals each reported on six other hospitals at the 1000-patient threshold (Figure 3C). The number
- 185 of hospitals in the covered set (achieving the threshold of >1000 received patients) was always
- 186 higher than the number of reporting hospitals (Figure 3D). In contrast, and by definition, any self-
- 187 reporting scheme reports only on exactly the numbers of hospitals included in the scheme.
- 188

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189 By selecting hospitals into the reporting set based on the number of patients they received from

- 190 other hospitals (labelled "receipt-based" in Figure 3D), the reach of the covered set could be
- substantially improved, with incidence estimated from >1000 patients in half the hospitals after
- 192 including just 16 hospitals in the reporting set. However, to estimate incidence in all hospitals, this
- 193 selection procedure still needed to include 101 hospitals in the reporting set.
- 194
- 195 A "greedy" algorithm significantly outperformed both the random and receipt-based additions to the
- 196 reporting set, increasing the covered set faster and providing the largest number of covered
- hospitals (with incidence estimated from >1000 patients) for any number of reporting hospitals. The
- difference between the greedy algorithm and the receipt-based selection was largest for the last 50
- 199 covered hospitals. Incidence could be estimated from >1000 patients in all hospitals after adding
- only 41 hospitals to the reporting set using the greedy algorithm (Figure 3D & 4A), while only 13
- 201 reporting hospitals were needed to survey 50% of all hospitals.
- 202

In the so-called "snowball" scenario, where hospitals start reporting if they are reported on, the number of reporting hospitals quickly expands. After the first hospital starts reporting its received

- 205 cases, its neighbours will join, followed by their neighbours, each time increasing the number of
- 206 received cases that are reported and the likelihood of other hospitals adding themselves to the
- 207 covered set (Figure 4B). For most randomly selected starting hospitals, this resulted in all hospitals
- 208 eventually being included in the reporting set. Only if the first hospital was small enough to not
- 209 receive >1000 patients from any particular hospital did the first step not result in the addition of
- 210 more hospitals to the reporting set (occurring with probability 19/155=0.12). For a group of nine
- 211 hospitals in the North, the snowball-addition stopped when the whole group was added, as the nine
- 212 hospitals combined did not receive >1000 patients from any other hospitals.
- 213

## 214 Discussion

- 215 To have the desired effect, incentives for hospitals to reduce their reported rates of AMR and other
- 216 hospital-transmitted organisms need to align with the hospitals' interests to reduce their numbers of
- colonised and infected patients. We show that this can be done by having hospitals report the
- 218 number of cases among the patients they admit who have previously been discharged from other
- 219 hospitals, as it separates the tasks of reporting and reducing incidence. In this way, hospitals report
- 220 on the AMR incidence in other hospitals, not on their own incidence, and as a result they
- themselves do not suffer potential consequences from their reports. Additionally, if the recipient
- hospital is then rewarded for any case they find, a clear incentive is constructed to find as many
- 223 cases as possible discharged from other hospitals, delivering a more reliable incidence estimate.
- 224
- The proposed surveillance system intrinsically increases the number of covered hospitals. First and foremost, by reporting cases admitted after previously being discharged from other hospitals, not all hospitals need to participate for it to be possible to estimate incidence for all hospitals. In fact, a
- selected subset of only 26% of English hospitals resulted in enough patients admitted to another
- 229 hospital within a year after discharge to estimate incidence in all hospitals in England. Even if
- 230 hospitals join the surveillance system (the reporting set) at random, incidences for all hospitals can
- 231 be obtained before all hospitals are reporting. The system therefore provides incidence estimates for
- 232 more hospitals than participate. Furthermore, because the reported incidence for a certain hospital
- will often be the result of the pooled reports sent in by several other hospitals, the final measured
- 234 incidence is less influenced by the screening rates of individual hospitals. The ranking of hospitals
- based on the agglomerated measurement can therefore be expected to be more robust than any
- 236 measurement derived from single hospitals.
- 237
- 238 The number of hospitals participating in such a surveillance scheme could easily increase if

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- 239 hospitals were compensated for cases they find among patients admitted after having been
- 240 discharged from another hospital, since there is no clear disadvantage to screening imported patients
- and reporting found cases. Subsequently, this effect may cause more hospitals to join: if a hospital's
- 242 incidence is reported by other hospitals, it may be inclined to start testing patients it admits after
- they have been discharged from other hospitals, if only to be able to compare incidences. Due to
- this snow-ball effect the system may not need to be mandatory, although a core group of
- 245 participating hospitals may be desirable.
- 246
- 247 If the goal of reporting incidence changes from purely gathering information to creating incentives
- for improving performance by penalising hospitals with high incidences, either financially or
- 249 reputationally, the proposed surveillance system still has value, because any repercussions
- associated with high incidence are incurred by a different hospital than the one that is screening
- 251 patients. However, exactly which cases might be counted when penalising hospitals needs to be
- carefully considered. To promote information sharing between hospitals, only newly discovered
- AMR-positive patients should be used to determine penalties, and not those patients that were previously screened and labelled as carriers, to prevent the punishment of hospitals that actively tr
- previously screened and labelled as carriers, to prevent the punishment of hospitals that actively try to share information about cases identified among their admitted patient population with other
- 255 to share information about cases identified among their admitted patient population with one 256 hospitals.
- 257

258 The proposed surveillance scheme exploits the structure of the hospital network, showing the added

259 value of regarding hospitals as interconnected by shared patients instead of completely independent

and isolated entities[8–12]. Previous studies have shown that patient sharing between hospitals

significantly correlate with rates of Carbapenemase-Producing Enterobacteriaceae (CPE)[13],

262 MRSA[14] or Clostridium difficile[15,16]. The influence of the hospital network formed by shared

263 patients on the spread of hospital-associated pathogens has also been used to design early warning

systems[16,17] or inform the distribution of resources for IPC[18], often reiterating the importance

265 of centrally located hospitals. We present a novel viewpoint on using these hospital networks, by

266 considering the interests of hospitals to report cases, thus actively using the shared patients to

- 267 combat the spread of these pathogens.
- 268
- 269 Limitations

270 The estimated incidence of a specific hospital measured by the reporting hospitals will not be

- 271 identical to incidence measured within the specific hospital itself, because the readmitted patients
- are a specific subset of the original patient population and more likely carriers. However, readmitted
- 273 populations will generally be broadly comparable between hospitals. Further, whilst this estimate
- may not precisely reflect the true incidence in a specific hospital, arguably neither does the self-
- 275 reported rate. Comparing estimated incidences for hospitals with vastly different function, such as

specialist hospitals, that have substantially different case-mix from the other hospitals, may need to

- be done carefully, for example using adjustment, as for standardised mortality rates.
- 278
- 279 We assumed that receiving hospitals are aware of patients' previous hospital stays upon admission,
- to identify those that need to be screened. However, this may not necessarily be the case, in
- 281 particular when the time since last discharge is relatively long. Reported incidences may therefore
- 282 be slightly lower, because some shared patients might be missed. Although this would lower the
- surveillance system's accuracy, the bias would be similar for all hospitals; in particular because
- 284 multiple hospitals can report on each covered hospital, any inaccuracies on the single reporting
- 285 hospital level will be averaged out.
- 286

We considered a cut-off for screening admissions of 1 year from previous discharge; in the general community, bacterial carriage may or may not persist over this period, making it harder to attribute bioRxiv preprint doi: https://doi.org/10.1101/535252; this version posted February 8, 2019. The copyright holder for this preprint (which was not certified by peer review) is the author/funder, who has granted bioRxiv a license to display the preprint in perpetuity. It is made available under a CC-BY-NC 4.0 International license.

289 colonisation status to the previous hospitalisation with confidence the longer a previous admission

290 was in the past. This is particularly problematic if levels of community transmission start to exceed

291 hospital-associated transmission. By shortening the cut-off time, the specificity of the surveillance

292 system will increase, at the cost of its sensitivity. However, by recording all colonised patients who

293 were previously admitted to another hospital, together with the time between admissions, it should

be possible to estimate the relative contribution of community transmission to the importation of

cases to all hospitals.

296

## 297 Conclusion

298 We propose a new system to estimate incidences of AMR and other hospital-transmitted micro-

299 organisms that does not rely on self-reporting, whereby instead surrounding hospitals report the

300 incidence within the patient population admitted to their hospital who have recently being

301 discharged from other hospitals. This decoupling of the hospital that is reporting from the hospital

302 reported on is vital for delivering reliable incidence estimates, as it takes away the incentive to stop

303 looking for cases by watching over the others. By reporting on other hospitals' incidence, the

304 surveillance scheme aligns financial and patient safety interests, encouraging hospitals to find and

- 305 report as many cases as possible, making the surveillance scheme more resilient against 'gaming'
- and thus delivering a more robust comparison between hospitals.
- 307

## 308 Supplementary information

- 309 Supplementary text: The game-theoretical implications of surveillance schemes
- 310
- 311 Supplementary table 1: Numbers of shared patients between hospitals, for cut-off time between
- admissions: one year, six months, three months, one month, and one week. Including list of hospital
- 313 codes and names.

## 314315 Abbreviations

- 316 AMR antimicrobial resistance
- 317 CPE carbapenemase-producing enterobacteriaceae
- 318 HES hospital episode statistics
- 319 IPC infection prevention and control
- 320 MRSA methicillin-resistant Staphylococcus aureus
- 321 NHS National Health Service
- 322

## 323 Authors' contributions

TD, ASW, and JVR designed the study. TD and TS performed the analysis. KLH provided data.

325 TD, TS, ASW, and JVR drafted the manuscript All authors revised the manuscript and gave final

- 326 approval for publication.
- 327

## 328 Funding

- 329 The research was funded by the National Institute for Health Research Health Protection Research
- 330 Unit (NIHR HPRU) in Healthcare Associated Infections and Antimicrobial Resistance at University
- of Oxford in partnership with PHE [grant number HPRU-2012-10041]. ASW, TEAP and DWC are
- 332 supported by the NIHR Oxford Biomedical Research Centre. TEAP and DWC are NIHR Senior
- 333 Investigators. The views expressed are those of the author(s) and not necessarily those of the NHS,
- the NIHR, the Department of Health or Public Health England.
- 335

## 336 Conflict of Interest

- 337 All authors declare no support from any organisation for the submitted work; NW has received
- 338 research grants from Wockhardt, Merck Sharp & Dohme Corp, Roche, Meiji Seika, Enigma

- 339 Diagnostics, Bio-Rad, Biomerieux, Accelerate, BD Diagnostics, Astrazeneca, Check points,
- 340 GlaxoSmithKline, Kalidex, Malinta, Momentum, Norgine, Rempex, Rotikan, Smith&Nephew,
- 341 Venato Rx Pharmaceuticals, and Basilea for research projects or contracted evaluations. There were
- 342 no other relationships or activities that could appear to have influenced the submitted work.
- 343

#### 344 Availability of data and materials

- 345 The patient admission data were not collected for this study specifically, and the authors do not own
- 346 the used datasets. Patient admission data from the English NHS HES (Hospital Episode Statistics.
- 347 Copyright© 2015. Re-used with the permission of the Health and Social Care Information Centre.
- 348 All rights reserved.) are available for researchers who meet the criteria for access to confidential
- 349 data from NHS digital (digital.nhs.uk; Formerly Health and Social Care Information Centre). All
- 350 other data needed to reproduce the analysis is provided in the supplementary information.
- 351

#### 352 **References**

- 1. Freeman T. Using performance indicators to improve health care quality in the public sector: a review of the literature. Heal Serv Manag Res. 2002;15:126–37.
- 2. Hibbard JH, Stockard J, Tusler M. Does publicizing hospital performance stimulate quality
   improvement efforts? Health Aff. 2003;22:84–94.

357 3. Maynard A, Bloor K. Will financial incentives and penalties improve hospital care? BMJ.
358 2010;340:c88.

- 359 4. Health Protection Agency. Quarterly Epidemiological Commentary: Mandatory MRSA, MSSA
- and E. coli bacteraemia, and C. difficile infection data (up to October December 2012). 2013;1–
  8.
- 362 5. Johnson AP, Davies J, Guy R, Abernethy J, Sheridan E, Pearson A, et al. Mandatory surveillance
- 363 of methicillin-resistant Staphylococcus aureus (MRSA) bacteraemia in England: the first 10 years. J
- 364 Antimicrob Chemother. 2012;67:802–9.
- 365 6. Wertheim HF, Melles DC, Vos MC, van Leeuwen W, van Belkum A, Verbrugh H a, et al. The
  366 role of nasal carriage in Staphylococcus aureus infections. Lancet Infect Dis. 2005;5:751–62.
- 367 7. Calfee DP, Durbin LJ, Germanson TP, Toney DM, Smith EB, Farr BM. Spread of methicillin-
- 368 resistant Staphylococcus aureus (MRSA) among household contacts of individuals with
- 369 nosocomially acquired MRSA. Infect Control Hosp Epidemiol. 2003;24:422–6.
- 8. Donker T, Wallinga J, Grundmann H. Patient referral patterns and the spread of hospital-acquired
   infections through national health care networks. PLoS Comput Biol. 2010;6:e1000715.
- 9. Iwashyna TJ, Christie JD, Moody J, Kahn JM, Asch DA. The structure of critical care transfer
  networks. Med Care. NIH Public Access; 2009;47:787–93.
- 374 10. Nekkab N, Astagneau P, Temime L, Crépey P. Spread of hospital-acquired infections: A
- 375 comparison of healthcare networks. PLOS Comput Biol. 2017;13:e1005666.
- 11. Huang SS, Avery TR, Song Y, Elkins KR, Nguyen CC, Nutter SK, et al. Quantifying
- interhospital patient sharing as a mechanism for infectious disease spread. Infect Control Hosp
   Epidemiol. 2010;31:1160–9.
- 12. Lee BY, McGlone SM, Song Y, Avery TR, Eubank S, Chang CC, et al. Social network analysis
- 380 of patient sharing among hospitals in Orange County, California. Am J Public Health.
- 381 2011;101:707–13.
- 382 13. Ray MJ, Lin MY, Weinstein RA, Trick WE. Spread of Carbapenem-Resistant

- 383 Enterobacteriaceae Among Illinois Healthcare Facilities: The Role of Patient Sharing. 2016;1–5.
- 14. Donker T, Wallinga J, Slack R, Grundmann H. Hospital networks and the dispersal of hospital acquired pathogens by patient transfer. PLoS One. 2012;7:e35002.
- 386 15. Simmering JE, Polgreen L a., Campbell DR, Cavanaugh JE, Polgreen PM. Hospital Transfer
- Network Structure as a Risk Factor for Clostridium difficile Infection. Infect Control Hosp
   Epidemiol. 2015;1–7.
- 389 16. Fernández-Gracia J, Onnela J-P, Barnett ML, Eguíluz VM, Christakis NA. Influence of a patient
- transfer network of US inpatient facilities on the incidence of nosocomial infections. Sci Rep.2017;7:2930.
- 392 17. Ciccolini M, Spoorenberg V, Geerlings SE, Prins JM, Grundmann H. Using an index-based
- approach to assess the population-level appropriateness of empirical antibiotic therapy. J
   Antimicrob Chemother. 2014;1–8.
- 395 18. Karkada UH, Adamic L a, Kahn JM, Iwashyna TJ. Limiting the spread of highly resistant
- 396 hospital-acquired microorganisms via critical care transfers: a simulation study. Intensive Care
- 397 Med. 2011;37:1633–40.

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#### 400 Figures



401

402 Figure 1) Schematic representation of the proposed surveillance system. A proportion of the

403 patients discharged from hospital 1 will be directly transferred or indirectly readmitted to hospitals

404 2-6. These shared patients may carry AMR acquired in hospital 1. By reporting these colonised

405 patients, as well as the total number of shared patients, hospitals 2-6 can estimate an AMR

406 incidence for hospital 1 without hospital 1 reporting.

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- 409 Figure 2) The English hospital network. A) The location of the included hospitals (dots), showing
- 410 the connections and connection weights based on patients shared between them (admitted to one
- 411 hospital having previously been discharged from another) (lines, darkness indicating the number of
- 412 shared patients). B) The distribution of connection weights between all hospitals. C) The
- 413 distribution of time between admissions, measured as days since previous discharge. D) The
- 414 distribution of lengths of stay, for all admissions (grey) and shared patients (blue).







417 Figure 3) A) The number of patients discharged from each hospital and subsequently admitted

418 elsewhere for different maximum periods between last discharge and next admission. If previous

discharges within a year are included, all hospitals discharge over 1000 patients who are

420 subsequently admitted elsewhere within a year. B) The number of hospitals that are covered by each

421 reporting hospital individually, as a function of the threshold number of received patients. C) The

422 number of hospitals that are covered by each reporting hospital individually, for a threshold of 1000

423 received patients (shown by red triangle in B). D) The number of hospitals covered as a function of

the number of reporting hospitals using self-reporting (black line) as well as the proposed

surveillance scheme with the reporting set determined by random assignment (grey), receipt-based

426 assignment (blue) and the greedy algorithm (blue).

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Figure 4) The geographical distribution of hospitals in the surveillance scheme. A) The minimal set of reporting hospitals needed to report on all hospitals, as found using the greedy algorithm. Green dots show the reporting set, grey dots the covered set and lines show the links over which patients

430 dots show the reporting set, grey dots the covered set and times show the times over which patients 431 previously discharged from other hospitals are included. B) The result of the snow-ball assumption

431 (a hospital will start reporting once it is reported on) as a function of the first hospital to join the

432 (a nospital will start reporting once it is reported on) as a function of the first nospital to join the

433 surveillance scheme. For the majority of hospitals (127/155), all other hospitals would join the

scheme were they the first hospital to start reporting (blue dots). However, a small group in the
 North region (9/155) will only report on hospitals in the same region (grey dots), while for small

North region (9/155) will only report on hospitals in the same region (grey dots), while for small
 number of hospitals (19/155) no others will join if they are the first in the surveillance system (red

437 dots), because they do not receive over 1000 patients per year from any other single hospital, and

437 dots), because they do not receive over 1000 patients per year from any other438 hence no other hospitals will therefore be reported on and join the scheme.

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