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Abstract

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Reference


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Variation in antibiotic use among and within different settings: a systematic review

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Objectives: Variation in antibiotic use may reflect inappropriate use. We aimed to systematically describe the variation in measures for antibiotic use among settings or providers. This study was conducted as part of the innovative medicines initiative (IMI)-funded international project DRIVE-AB.

Methods: We searched for studies published in MEDLINE from January 2004 to January 2015 reporting variation in measures for systemic antibiotic use (e.g. DDDs) in inpatient and outpatient settings. The ratio between a study’s reported maximum and minimum values of a given measure [maximum:minimum ratio (MMR)] was calculated as a measure of variation. Similar measures were grouped into categories and when possible the overall median ratio and IQR were calculated.

Results: One hundred and forty-three studies were included, of which 85 (59.4%) were conducted in Europe and 12 (8.4%) in low- to middle-income countries. Most studies described the variation in the quantity of antibiotic use in the inpatient setting (81/143, 56.6%), especially among hospitals (41/81, 50.6%). The most frequent measure was DDDs with different denominators, reported in 23/81 (28.4%) inpatient studies and in 28/62 (45.2%) outpatient studies. For this measure, we found a median MMR of 3.7 (IQR 2.6–5.0) in 4 studies reporting antibiotic use in ICUs in DDDs/1000 patient-days and a median MMR of 2.3 (IQR 1.5–3.2) in 18 studies reporting outpatient antibiotic use in DDDs/1000 inhabitant-days. Substantial variation was also identified in other measures.

Conclusions: Our review confirms the large variation in antibiotic use even across similar settings and providers. Data from low- and middle-income countries are under-represented. Further studies should try to better elucidate reasons for the observed variation to facilitate interventions that reduce unwarranted practice variation. In addition, the heterogeneity of reported measures clearly shows that there is need for standardization.

Introduction

Variation in healthcare delivery among different geographical areas, healthcare facilities and individual providers is a nearly ubiquitous finding that can only be partly explained by differences in patient characteristics or disease epidemiology.1,2 The importance of systematically studying the extent of this variation and its underlying causes in order to improve quality and resource utilization was first recognized in the 1970s by Wennberg and Gittelsohn,3 when they analysed ‘small area variations’ of hospitalization rates and surgical procedures in the US state of Vermont. With the continuing emergence and spread of MDR organisms across the globe in recent years, there has been increasing interest in medical practice variation regarding antibiotics since inappropriate use is one of the key drivers of antimicrobial resistance.4–8 Indeed, important variation in antibiotic use has been described among countries, hospitals and physicians that share many similarities in their patient populations and economic, geographical
Eligibility criteria

We conducted a systematic review of the published literature to identify English language studies that describe the naturally observed variation (i.e. the variation occurring outside the context of a specific interventional study) in measures of systemic antibiotic use for treatment and prophylaxis within and among different settings (e.g. countries, hospitals, hospital units) and providers (e.g. general practitioners, paediatricians, physicians with different specialities) at a given point in time. We included studies describing variation in both paediatric and adult populations. Only studies describing variation among a minimum number of providers or settings were included (Table 1). We predefined different limits for larger entities (such as hospitals and countries) and individual providers (Table 1). The cut-offs were chosen for pragmatic reasons, since we felt that otherwise the number of eligible studies would be too vast without offering much information about variation due to the small number of entities.

<table>
<thead>
<tr>
<th>Setting</th>
<th>Unit/hospital/region/country level</th>
<th>Provider level</th>
</tr>
</thead>
<tbody>
<tr>
<td>Inpatient</td>
<td>Data from ≥5 hospitals irrespective of their size OR Data from ≥5 identical units (e.g. ICUs, haematology wards etc.) from ≥5 hospitals irrespective of their size OR Data from ≥5 units/wards in the same hospital irrespective of their size</td>
<td>≥20 providers in the same hospital</td>
</tr>
<tr>
<td>Outpatient</td>
<td>≥2 countries, regions or districts (same or different country) OR ≥5 clinics/primary health care facilities</td>
<td>≥50 providers in the same geographical area</td>
</tr>
</tbody>
</table>

The cut-offs were chosen for pragmatic reasons, since we felt that otherwise the number of eligible studies would be too vast without offering much information about variation due to the small number of entities.

Methods

This systematic review is reported following the PRISMA statement.19

Eligibility criteria

We conducted a systematic review of the published literature to identify English language studies that describe the naturally observed variation (i.e. the variation occurring outside the context of a specific interventional study) in measures of systemic antibiotic use for treatment and prophylaxis within and among different settings (e.g. countries, hospitals, hospital units) and providers (e.g. general practitioners, paediatricians, physicians with different specialities) at a given point in time. We included studies describing variation in both paediatric and adult populations. Only studies describing variation among a minimum number of providers or settings were included (Table 1). We predefined different limits for larger entities (such as hospitals and countries) and individual providers (Table 1). The cut-offs were chosen for pragmatic reasons, since we felt that otherwise the number of eligible studies would be too vast without offering much additional information about variation due to the small number of entities.

We allowed the inclusion of ESAC/ESAC-net (European Surveillance of Antimicrobial Consumption Network) studies reporting data from the same year(s) only when the number of countries participating in the studies was different or the described measures were different.

We tried to group similar measures into categories reflecting:

- The quantity of antibiotic use [e.g. DDDs, days of therapy (DOT), length of therapy (LOT) with different denominators or percentage of treated patients].
- Prescribing strategies (e.g. percentage of delayed prescriptions, percentage of antibiotics prescribed as empirical treatment).
- Compliance with guidelines (percentage of appropriate or compliant prescriptions, percentage of patients treated with antibiotics within a given timeframe).
- Process and structural measures for antibiotic stewardship policies (e.g. percentage of prescriptions documented in the medical file, presence of antibiotic stewardship guidelines).
- Antibiotic use for medical or surgical prophylaxis (e.g. percentage of patients receiving surgical prophylaxis for >24 h).

We excluded the following studies:

- Studies describing exclusively the use of antivirals, antifungals, antimycobacterials, antiparasitic drugs and topical antibiotics.
- Studies only describing variation of antibiotic use over time within the same setting.
- Studies only describing variation in outcomes associated with antibiotic use (e.g. variation in rates of Clostridium difficile infection).
- Studies focusing on the variation in healthcare professionals’ views, beliefs, attitudes and knowledge.
- Studies describing self-reported (by patients, caregivers and physicians) behaviours regarding antibiotic use.
- Studies whose full text could not be retrieved from any of the libraries of the participating centres (eight different catalogues).
- Studies that presented data only graphically (no efforts were made to contact authors).
- Interventional studies without extractable pre-intervention data.
- Systematic reviews and meta-analyses and studies not reporting original data (narrative reviews, opinion pieces etc.). Their reference lists were, however, screened to identify potentially eligible studies.

Search and information sources

We searched the MEDLINE database using the PubMed interface using a combination of search terms for the concepts (i) ‘antibiotics’, (ii) ‘quality’ and ‘quantity of use’ and (iii) ‘variation’ (for the detailed search strategy see
Table S1, available as Supplementary data available at JAC Online). Owing to the large number of potentially eligible studies (>6000) we used the PubMed filters for species (‘humans’) and language (‘English’). Since we were mostly interested in recent findings, we restricted the timeframe to articles whose last year of data collection was 2004 (included) or later with a publication date between 1 January 2004 and 15 January 2015.

Study selection and data collection process

All the steps of this systematic review were carried out using the Distiller SR© software (Evidence Partners, Ottawa, Canada). Duplicates were removed before title and abstract screening using the algorithm provided by Distiller SR©. One reviewer (V. Z.) screened all titles and abstracts. A second reviewer (B. H.) independently screened a random subset of 700 abstracts (13%). For 3/700 (0.4%) references screened by both reviewers there was disagreement regarding inclusion/exclusion. All three references were later excluded at full-text screening level.

Full-text assessments were performed by one reviewer (V. Z.) and, in case of uncertainty, discussed with another investigator (B. H.) until a consensus was reached. Data extraction was performed by one reviewer (V. Z.) using a standardized data extraction form; any uncertainty about extracted data was discussed with another investigator (B. H.).

Data regarding authors, setting (inpatients or outpatients), country/region, last year of data collection, study design, level of variation (providers, units, hospitals etc.), number and characteristics of participants, data source, category of the numerator and denominator of the measure, full description of the measure, mean or median (as available in the text) and measure of variability (maximum–minimum range, IQR or SD as presented in the text) were collected (for definitions see Table S2). We extracted data for overall antibiotic use and if the study also reported variation for specific antibiotic classes we extracted data only for β-lactams, quinolones and macrolides (except if the study described variation of only specific antibiotic classes but no overall antibiotic use) since we felt that extracting data for all individual classes would go beyond the scope of this review. Since we mainly included observational studies and the aim of this review was to describe variation (without causal inferences) we decided not to perform risk of bias assessment for the included studies.

Synthesis of results and summary measures

Using the search strategy (Table S1), we identified 5204 studies. After title and abstract screening 628 studies were retained for full-text review; of these 143 met the inclusion criteria (Figure 1). Selected study characteristics are presented in Table 2. Detailed information for all 143 studies is available in Tables S5 to S8. The results of our search are presented by setting (inpatients versus outpatients) and country income level (high versus low to middle income using the World Bank 2015 classification20); for high-income countries we further differentiated between European and North
American studies since these were the most represented regions. Data regarding measures for antibiotic use in paediatric patients are presented as a separate category. Measures were extracted from each study and a measure’s category was attributed to each of them. The complete list of measure categories is reported in Table S3.

Given our broad search strategy, we identified a wide range of different antibiotic use measures, often applying to specific medical conditions (e.g. otitis media, urinary tract infections or community-acquired pneumonia) or settings (e.g. ICUs, neonatal ICUs, emergency departments). In addition, most studies presented only aggregated data, making it challenging to summarize variation in a standardized way. In the absence of better alternatives, we decided, whenever possible, to report the extent of variation in a given antibiotic use measure for each study by calculating the ratio between the reported maximum and minimum value of the measure (MMR). For example, in a study describing variation in overall antibiotic use among hospitals in DDDs we divided the maximum value by the minimum value of DDDs. This ratio was then used to calculate the median and IQR ratio whenever measures of the same category could be grouped together.

Results

Overall a total of 44 unique measures grouped in five categories were identified (Table S3). For the purpose of this review the term ‘unique measures’ is used to indicate distinctive measures, meaning measures different from one another with respect to their numerator (e.g. DDDs, DOT, percentage of treated patients). Measures that have the same numerator but different denominators, e.g. DDDs per inpatient day or per admission, were not considered unique measures for the purpose of this review because the numerator is the same. An overview of the frequency of the measures in the different categories both in the in- and outpatient settings is presented in Table S4.

In the context of this review we identified 10 unique measures for quantity, 5 for prescribing strategies, 8 for compliance with guidelines, 18 for process and structural measures of antibiotic stewardship policies and 3 for antibiotic prophylaxis.

Inpatient setting (81 studies)

High-income countries

Overall, 75 studies (75/143, 52.4%) met inclusion criteria for this setting: 47 from Europe, 17 from the USA and Canada, 3 from Australia, 1 from Israel, 1 from Japan and 6 from more than one WHO region (see Table S2 for the definition of WHO regions). Thirty-eight unique measures were extracted for this setting with sometimes different studies reporting identical measures. Overall, 169 ‘non-unique’ measures reporting data about variation (also called ‘variation data’ in this review) were represented for this setting. The most frequent measures concerned ‘quantity’ of antibiotic use, with 91 variation data extracted from 58 studies. The three most frequent unique measures (out of the 10 belonging to this category) were DDDs (reported 34 times for overall use or for specific antibiotic classes and medical conditions and with different denominators), followed by ‘percentage of treated patients’ (reported 31 times in different settings and for different health conditions) and DOT (reported 13 times both for overall and specific antibiotic classes use and for specific medical indications and with different denominators).
Table 3. Most frequent measures, Europe and North America: inpatient setting (64 studies)

<table>
<thead>
<tr>
<th>Measure</th>
<th>Number of studies (%)</th>
<th>MMR (IQR) (if applicable)</th>
<th>Comments</th>
</tr>
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<tbody>
<tr>
<td><strong>Europe</strong></td>
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</tbody>
</table>
| percentage of patients treated with antibiotics (any condition)         | 19/47 (40.4)          | Hospitals (5 studies29–33); median MMR 2.1 (1.8–2.4) | 29PPS in 30 Finnish hospitals in 2005  
20PPS to determine the prevalence of antibiotic use for hospital-acquired infections in 393 French non-teaching hospitals in 2001 and 2006  
21PPS of nosocomial infections in 41 Dutch hospitals in 2007 and 2008  
22PPS to determine the prevalence and the appropriateness of antimicrobial use in 19 Dutch hospitals in 2008 and 2009  
23ESAC PPS of antibacterial use in 20 European hospitals in 2006  
24Cross-sectional study in 18 nursing homes in Franche-Comté (France) on residents receiving antibiotics on the study day in 2012  
25PPS in 30 nursing homes in Northern Ireland in 2011  
26PPS in 85 nursing homes in 15 European countries and two UK administrations in 2009  
27Two PPSs in 30 nursing homes in Northern Ireland in 2009  
28PPS in 9 Finnish nursing homes in 2010  
29PPS in 323 nursing homes in 21 European countries in 2009  
30Cross-sectional study in Franche-Comté (France) on residents receiving antibiotics on the study day in 2012  
31PPS in 30 nursing homes in Northern Ireland in 2011  
32PPS to determine the prevalence and the appropriateness of antimicrobial use in 19 Dutch hospitals in 2008 and 2009  
33ESAC PPS of antibacterial use in 20 European hospitals in 2006  
34Cross-sectional study in 18 nursing homes in Franche-Comté (France) on residents receiving antibiotics on the study day in 2012  
35PPS in 30 nursing homes in Northern Ireland in 2011  
36PPS in 85 nursing homes in 15 European countries and two UK administrations in 2009  
37Two PPSs in 30 nursing homes in Northern Ireland in 2009  
38PPS in 9 Finnish nursing homes in 2010  
39PPS in 323 nursing homes in 21 European countries in 2009  |
| total DDDs/1000 patient-days                                            | 16/47 (34.0)          | Hospitals (6 studies40–45); median MMR 4.2 (2.3–8.1) | 40Cohort study of inpatient antibiotic use in acute hospitals in England analysed over 5 years (2008–13)  
41Ecological study in French healthcare facilities  
42Study on antibiotic consumption in 57 Swiss hospitals  
43–45Studies on antibiotic consumption in French hospitals |
| all measures related to antibiotic stewardship                          | 7/4749–55 (14.9)      | NA                         | Examples of measures: presence of a surveillance system for antibiotic use or presence of an antibiotic stewardship programme for LTCFs  
50Appropriate antibiotic prescriptions for urinary tract infections in emergency departments of 10 hospitals from different Spanish regions |
| percentage of appropriate prescriptions                                 | 1/47 (2.1)            | Emergency departments (1 study56); MMR 1.5         | 56In this study, antibiotic use was measured using number of use-days/100 patient-days during a 7 day period. The study reported the prevalence of patients receiving at least two antimicrobials during the study day in 30 Finnish hospitals |
| antibiotic days/patient-days                                            | 2/47 (4.2)            | Hospitals (1 study55); MMR 2.1 | 55In this study, antibiotic use was measured using number of use-days/100 patient-days during a 7 day period. The study reported the prevalence of patients receiving at least two antimicrobials during the study day in 30 Finnish hospitals |
| number of antibiotics accounting for 75% of total consumption          | 1/47 (2.1)            | Hospitals (1 study57); MMR 2.1 | 57Drug utilization 75% (DU75%) in 17 European hospitals: results from the ESAC-2 Hospital Care Sub Project |
| percentage of combination therapy                                       | 1/47 (2.1)            | Countries (1 study58); MMR 1.2 | 58European Surveillance of Antimicrobial Consumption (ESAC): value of a PPS of antimicrobial use across Europe. The use of combination therapy was related to hospital type, with teaching and tertiary hospitals having a significantly higher use of combination therapy |
| percentage of surgical prophylaxis >24 h                                | 1/47 (2.1)            | Countries (1 study59); MMR 3.1 | 59Prolonged perioperative surgical prophylaxis within European hospitals. For the purpose of the study data were extracted from the ECDC PPS 2011–12 report |
| percentage of patients receiving prophylaxis (medical or surgical)     | 4/47 (8.5)            | LTCFs (2 studies): MMR 22 (average60), MMR 55 (for UTI61) | 56Variation was described among 44 Norwegian LTCFs. 10% of residents on the day of the survey were receiving antibiotics for infection prevention and 6% for infection treatment. The indication for prophylaxis was UTI in all but 1 case  
57Timely prophylaxis before surgery (within 1 h of procedure)  
58Percentage of patients receiving prophylaxis before cystoscopy across hospitals in different countries (data from the Global Prevalence Study on Infections in urology) |

Continued
Europe

Forty-seven studies (47/143, 32.9%), including 9 ESAC studies, were included for the inpatient setting, reporting data from 13 different European countries. We identified 18 variation data specifically addressing the paediatric population and 4 variation data concerning antibiotic prophylaxis.

Included studies described variation either among acute care hospitals (18/47, 38.3%), long-term care facilities (LTCFs) (7/47, 14.9%), neonatal ICUs (2/47, 4.3%), adult ICUs (6/47, 12.8%), different hospital units (10/47, 21.3%) or different countries (4/47, 8.5%). No study described variation among providers (e.g. physicians with different specialties). Concerning the nine ESAC studies, almost all (8/9, 88.9%) addressed antibiotic use for treatment in the adult population, describing variation among hospitals (3/9, 33.3%), units (2/9, 22.2%), LTCFs (2/9, 22.2%) or countries (2/9, 22.2%). Both among units and among LTCFs, the most frequently reported measure was overall percentage of patients treated with an antibiotic (only one study specifically addressed nosocomial infections at the unit level).

We identified 58 variation data for this setting for the category ‘quantity’ of antibiotic use from 37 studies (Table S5) for the complete list of measures with calculated levels of variation (MMR).

MMRs for the most frequently reported measures referring to the same population, medical condition and setting are presented in Table 3. The highest MMR for this setting was described among LTCFs (six studies) for the measure ‘percentage of patients treated with antibiotics for any condition’, with a median MMR of 13.3 (IQR 8.9–16.5). The same measure presented a median MMR of 2.1 (IQR 1.8–2.4) among hospitals (five studies). We also found a median MMR of 4.2 (IQR 2.3–8.1) among hospitals (six studies) and of 3.7 (IQR 2.6–5.0) among ICUs (four studies) for the measure DDDs/1000 patient-days.

North America

We found 17 North American studies that fulfilled inclusion criteria for the inpatient setting: 14 from the USA and 3 from Canada; 53% (9/17) addressed the paediatric population and 2 described variation in measures for antibiotic use in LTCFs. Three studies included measures for antibiotic prophylaxis. One was an interventional study. The four most frequent unique measures described for this setting are reported in Table 3. Among the paediatric studies the MMR for the measure ‘percentage of febrile neonates treated

Table 3. Continued

<table>
<thead>
<tr>
<th>Measure</th>
<th>Number of studies (%)</th>
<th>MMR (IQR) (if applicable)</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>percentage of patients whose antibiotic prophylaxis was stopped &lt; 24 h after surgery</td>
<td>1/47 (2.1)</td>
<td>Departments (1 study): MMR 1.2</td>
<td>62*The measure specifically referred to duration of surgical prophylaxis &lt; 24 h in Scottish acute care hospitals</td>
</tr>
<tr>
<td>North America</td>
<td>percentage of patients treated with antibiotics</td>
<td>2/17 (11.8)</td>
<td>Departments (1 study addressing adults): MMR 26.6; LTCFs (1 study): MMR 4.5</td>
</tr>
<tr>
<td>DOT/antibiotic days</td>
<td>6/17 (35.3)</td>
<td>Hospitals (1 study): MMR 1.8</td>
<td>66*This study examined the relationship between nursing home prescriber adherence to the Loeb minimum criteria and antibiotic prescribing rates, overall and for each of three specific conditions (urinary tract infections, respiratory infections, and skin and soft tissue infections)</td>
</tr>
<tr>
<td>prophylaxis use before surgery</td>
<td>3/17 (17.6)</td>
<td>Hospitals (3 studies): Adults median MMR 4.5</td>
<td>67*Variation was measured among 70 US academic centres</td>
</tr>
<tr>
<td>DDDs/10000 patient-days</td>
<td>1/17 (5.9)</td>
<td>Hospitals (1 study): MMR 2.5</td>
<td>68*Interventional study examining the effect of antibiotic stewardship programme (ASP)-based strategies (all including a component of audit and feedback) on antibiotic consumption of target antibiotics (piperacillin/tazobactam, fluoroquinolones, or cefepime) Data refer to the baseline data in the intervention group (non-target antibiotics)</td>
</tr>
<tr>
<td>percentage of patients whose antibiotic prophylaxis was stopped &lt; 24 h after surgery</td>
<td></td>
<td>Hospitals (1 study): MMR 4.5</td>
<td>62*The measure specifically referred to duration of prophylaxis &lt; 24 h across 295 US hospital groups (a hospital group comprises all hospitals sharing identical categories for location by state, teaching status, bed size and urban/rural location)</td>
</tr>
</tbody>
</table>

PPS, point prevalence study.

*The Loeb minimum criteria, developed by a 2001 consensus conference, are minimum standards for initiation of antibiotics in long-term care settings, intended to reduce inappropriate prescribing.
with antibiotics' among departments was 1.321 and 1.522 in two studies, respectively.

An MMR of 2.5 was described in one study describing the same measure in children with bronchiolitis presenting at the emergency department. As in Europe, the MMR among LTCFs for the percentage of patients receiving antibiotics was high (4.5 in one study).

Low- to middle-income countries

We found three different measures described for this setting (Table 4), two belonging to the category ‘quantity’ (DDDs and percentage of treated patients) and one belonging to the category ‘compliance with guidelines’ (percentage of appropriate/inappropriate prescription).

Six studies could be included for this setting (three of which were from the Western Pacific WHO region); one was an international study that included countries from more than one WHO region and the others were from four different countries. No study included data collected after 2011. Three studies addressed the paediatric population. Variation was mainly described among hospitals (four studies, 66.7%) with an MMR of 2.1 (IQR 2–2.2) for the measure ‘percentage of patients treated with antibiotics’ (three studies) and an MMR of 5.1 for the measure DDDs/100 bed-days (one study). The complete list of included studies for this setting is presented in Table S6.

Outpatient setting (62 studies)

High-income countries

Overall, 56 studies (56/143, 39.2%) met the inclusion criteria for this setting: 38 from Europe, 10 from the USA and Canada, 1 each from Bahrain, Israel, Saudi Arabia and South Korea and 4 from more than one WHO region. Overall, 110 variation data were identified (Table S7).

As for the inpatient setting, the category with the highest number of included measures was ‘quantity’ of antibiotic use, with 53 studies reporting 91 variation data. The three most frequent measures were: (i) DDDs (34 variation data in 28 studies); (ii) percentage of antibiotic prescriptions for different medical conditions, populations (e.g. children, adults or elderly people) and sometimes for specific antibiotic classes (18 variation data in 10 studies); and (iii) percentage of treated patients (17 variation data in 12 studies).

Europe

We included 38 studies (38/56, 67.8%) for this setting, including 15 ESAC studies. No study reported data collected after 2011. More than half were international studies (20/38, 52.6%); the rest were single-country studies from nine different countries. Italy was the country with the highest number of studies (6/38, 15.8%) followed by France (4/38, 10.5%).

Most of the studies reported variation among countries (15/38, 39.5%) or geographical areas within the same country (e.g. regions, provinces, districts) (7/38, 18.4%). Ten studies (26.3%) reported variation among providers and six among outpatient clinics. Overall, most studies (36/38, 94.7%) included measures for the category ‘quantity’ of antibiotic use, with the three most represented measures being: (i) DDDs (reported in 21 studies); (ii) percentage of treated patients (reported in 10 studies); and (iii) percentage of antibiotic prescriptions (reported in 9 studies). The second category in terms of frequency of measures was ‘prescribing strategies’ [e.g. percentage of delayed prescriptions for lower respiratory tract infections (LRTIs), days of delay before taking the antibiotic for LRTIs, percentage of patients treated with specific antibiotic classes for respiratory tract infections, and percentage of antibiotics administered through parenteral route] with 10 unique measures included followed by ‘compliance with guidelines’ [e.g. percentage of guideline-compliant prescriptions for LRTIs] with 6 unique measures included. No study reporting measures for the category ‘process and structural measures for antibiotic stewardship policies’ was identified for this setting.

In Table 5 we report the level of variation for the most frequent measures: DDDs/1000 inhabitant-days, which showed a median MMR of 3.2 (IQR 3.0–3.5) among countries (eight studies) and a median MMR of 1.5 (IQR 1.4–1.5) among geographical areas (three studies). For the measure ‘percentage of treated patients’ we found an MMR of 1.4 in two studies reporting variation among geographical areas in the same country.

North America

Ten studies (10/62, 16.1% of all studies included for the outpatient setting) describing variation in outpatient antibiotic use in North America were identified: 6 from Canada, 3 from the USA and 1 that included data from both countries. Two studies (20%) included measures addressing the paediatric population. No study described measures for antibiotic prophylaxis.

Nine studies included measures for the category ‘quantity’ of antibiotic use and one study included one measure for the category ‘process and structural measures for antibiotic stewardship policies’. No study described measures for the remaining three categories (‘compliance with guidelines’, ‘prescribing strategies’ and ‘antibiotic prophylaxis’).
The most frequent measure reported for this setting (Table 5) was DDDs/1000 inhabitant-days (six studies), with an MMR of 1.9 among geographical areas (two studies). An MMR of 1.4 was described also for the measure ‘percentage of treated patients’, again among geographical areas in the same country (one study).

### Table 5. Outpatient setting: Europe and North America (48 studies)

<table>
<thead>
<tr>
<th>Measure</th>
<th>Number of studies (%)</th>
<th>MMR (maximum/minimum ratio) (IQR)</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Europe</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>DDDs/1000 inhabitant-days</td>
<td>19/38 (50.0)</td>
<td>Countries (8 studies⁶⁻¹⁰,⁷²⁻⁷⁷):</td>
<td>No study included data collected after 2009 8 were ESAC studies</td>
</tr>
<tr>
<td></td>
<td></td>
<td>median MMR 3.2 (3.0–3.5)</td>
<td></td>
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<tr>
<td></td>
<td></td>
<td>Geographical areas (3 studies⁷⁸⁻⁸⁰):</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>median MMR 1.5 (1.4–1.5)</td>
<td></td>
</tr>
<tr>
<td>percentage of treated patients</td>
<td>12/38 (31.6)</td>
<td>Geographical areas (2 studies⁶¹,⁸²):</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>MMR 1.4</td>
<td></td>
</tr>
<tr>
<td>percentage of total antibiotic use duration of therapy</td>
<td>1/38 (2.6)</td>
<td>Providers (1 study⁸³)</td>
<td></td>
</tr>
<tr>
<td>percentage of compliant prescriptions</td>
<td>1/38 (2.6)</td>
<td>Clinics (1 study⁸⁵):</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>MMR 6.6</td>
<td></td>
</tr>
<tr>
<td>percentage of delayed prescriptions</td>
<td>1/38 (2.6)</td>
<td>Clinics (1 study⁸⁵):</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>MMR 16.5</td>
<td></td>
</tr>
<tr>
<td><strong>North America</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>DDDs/1000 inhabitant-days</td>
<td>6/10 (60)</td>
<td>Geographical areas (2 studies⁷⁷,⁸⁸):</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>MMR 1.9</td>
<td></td>
</tr>
<tr>
<td>prescriptions</td>
<td>2/10 (20.0)</td>
<td>Geographical areas (1 study⁸⁵):</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>MMR 1.4</td>
<td></td>
</tr>
<tr>
<td>percentage of treated patients</td>
<td>2/10 (20.0)</td>
<td>Providers (1 study⁸⁹):</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>MMR 2.3</td>
<td></td>
</tr>
<tr>
<td>percentage of non-compliant prescriptions</td>
<td>1/10 (10.0)</td>
<td>Clinics (1 study⁹¹):</td>
<td></td>
</tr>
<tr>
<td>number of antibiotics whose prescription is restricted and requires specific information confirming the diagnosis in order for the patient to be reimbursed by the health system</td>
<td>1/10 (10.0)</td>
<td>Geographical areas (1 study⁹²):</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>MMR 15</td>
<td></td>
</tr>
</tbody>
</table>

The most frequent measure reported for this setting (Table 5) was DDDs/1000 inhabitant-days (six studies), with an MMR of 1.9 among geographical areas (two studies). An MMR of 1.4 was described also for the measure ‘percentage of treated patients’, again among geographical areas in the same country (one study).

### Low- to middle-income countries

Six studies fulfilled the inclusion criteria (9.7% of all studies included for the outpatient setting). One was an international study and the rest were single-country studies from three different countries (China, Iran and India). One interventional study was...
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Table 6. Low- to middle-income countries: outpatient setting (6 studies)

<table>
<thead>
<tr>
<th>Measure</th>
<th>Number of studies (%)</th>
<th>MMR, median (IQR) (if applicable)</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>Percentage of treated patients</td>
<td>3/6 (50)</td>
<td>Providers (1 study\textsuperscript{93}): 20.5</td>
<td>\textsuperscript{93} Antibiotic use among GPs and specialist physicians in Iran (this survey was conducted on a total of almost 8 million prescriptions)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Clinics (1 study\textsuperscript{96}): 1.4</td>
<td>\textsuperscript{96} Surveillance of antibiotic encounters carried out using a repeated cross-sectional design for 2 years in Vellore, South India. Variation was described among 30 health facilities</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Geographical areas (1 study\textsuperscript{97}): 1.4</td>
<td>\textsuperscript{97} Antibiotic use in rural areas of 10 Chinese provinces</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Countries (1 study\textsuperscript{98}): 2.4</td>
<td>\textsuperscript{98} Antibiotic use in eight Latin American countries (we considered only data from 2007)</td>
</tr>
<tr>
<td>DDDs/1000 inhabitant-days</td>
<td>2/6 (33.3)</td>
<td>(one of the studies concerned the paediatric population)</td>
<td></td>
</tr>
<tr>
<td>Percentage of antibiotics per prescription</td>
<td>1/6 (16.7)</td>
<td>Providers (1 study\textsuperscript{99}): 2.2</td>
<td>\textsuperscript{99} Cluster randomized controlled trial of 159 GPs working in 6 cities, in 2 regions in East Azerbaijan in Iran. The cities were matched and randomly divided into an intervention arm, for an outcome-based education on rational prescribing, and a control arm for a traditional CME programme on the same topic. GPs’ prescribing behaviour was assessed 9 months before and 3 months after the CME programmes</td>
</tr>
</tbody>
</table>

CME, continuing medical education; GPs, general practitioners.

Discussion

The most important finding of this review was the large variation in measures of antibiotic use even across similar settings and providers. This finding was confirmed also for specific medical conditions and populations both in high- and low- to middle-income countries. A second key finding is the large heterogeneity of reported measures (even without taking into account differences in data sources; e.g. antibiotic dispensing versus administration data), clearly indicating the need for standardization.

Most of the findings of this review concerned evaluation of the quantity of antibiotic consumption, and variation was observed whatever the type of measure used and regardless of the setting. For example, for the European inpatient setting we found high variation in DDDs/1000 patient-days with a median MMR of 4.2 for hospitals (six studies) and 3.7 for ICUs (four studies). In low- to middle-income countries the same measure showed an MMR of 5.1 (one study) among hospitals. Variation was also observed for antibiotic prescribing strategies, guideline compliance and for existence of antimicrobial stewardship policies across healthcare facilities. In most of the cases measures referred to antibiotic use for treatment in the adult population, whereas studies including antibiotic use for prophylaxis or addressing the paediatric population were a minority. It was not surprising to see that most of the variation was described among providers (two studies), countries or smaller geographical areas within the same country (two studies) or outpatient clinics (two studies). We identified three different measures for this setting (Table 6), with the highest MMR (20.5) described among providers (one study) for the measure ‘percentage of treated patients’ in a study from Iran. The complete list of included studies for this setting is presented in Table S8.
information retrieved from the literature came from observational retrospective studies from high-income countries, especially from Europe, and that data from resource-limited settings were under-represented in the literature.23

Despite the fact that our search was performed at the beginning of 2015, most data were from before 2012, most likely reflecting the delay between data collection, data analysis and final publication. The great variety of measures made it difficult to summarize and present the observed variation. The ratio between maximum and minimum values is certainly a suboptimal measure of variation since it is heavily influenced by outliers and the characteristics of the measure (e.g. MMRs will be inherently different for measures with an absolute upper and lower boundary, such as those expressed in percentages, and measures that only have an absolute lower boundary, such as DDDs).

Inferences on the causes of the described variation were beyond the scope of this review. Indeed, few of the examined studies offered clear evidence for the reasons behind the observed variation. It seems clear, however, that much of the observed variation is ‘unwarranted’, in the sense that it is unlikely to be driven by differences in the epidemiology of infectious diseases or patient characteristics but could rather be explained by one or more of the seven determinants of healthcare professionals’ behaviours described by Flottorp et al.24 and that could also be applied to antibiotic prescribing practices. These seven domains are: (i) guideline factors (e.g. guideline characteristics or presence of contradictory guidelines); (ii) individual prescriber preferences (which could be influenced by many factors, such as lack of agreement with specific guidelines, lack of motivation or inertia of previous practices etc.); (iii) patient factors (e.g. patient’s expectations, inability to reconcile patient’s preferences with guideline recommendations); (iv) professional interactions (e.g. leadership, key individuals, team processes); (v) incentives and resources (e.g. economic incentives, technical knowledge, organizational size); (vi) capacity for organizational change (e.g. planning, engaging, executing and evaluating); and (vii) social, political and legal medical norms (e.g. legislation or regulations, priority on societal agenda, corruption, political stability etc.). The impact of combinations of these domains on antibiotic use has been previously described in both the inpatient and the outpatient setting.25–28

In particular in the inpatient setting, differences in ‘culture’ (country level, e.g. ideas about health, causes of disease, labelling of illness, coping strategies and ‘treatment modalities’ differ across countries), ‘context’ (hospital level, influenced by organizational policies, multi-professional care-delivery system) and ‘behaviour’ (professional level) have all been described.28 In the outpatient setting, socio-cultural differences as well as specific regulatory practices have been associated with varying levels of prescriptions for frequent medical conditions, such as LRTIs.27

Strengths of this review include a broad search strategy and the inclusion of studies from a variety of different settings. There are, however, also several limitations to our work. We only searched the MEDLINE database over a 10-year period and we did not explore other sources of information, such as relevant surveillance websites or single-country data. Another limiting factor is that we did not perform a quality assessment of included studies and we did not explore the relationship between extent of variation in measures and outcomes of antibiotic use, e.g. resistance, costs or rates of C. difficile infections. Although we included a large variety of measures described in the literature, we did not address possible benchmarks for the measures and we did not try to explore reasons for the observed variation as most studies did not provide information regarding this issue. We did not address the relevance of the measures for appropriate use. This is addressed in other work of the DRIVE-AB consortium.14–18

Conclusions
At a time when major stakeholders are trying to find effective solutions to the problem of antibiotic resistance, the large variation in measures of antibiotic use (even across similar settings/providers or for similar clinical conditions) remains poorly understood and can only be partly explained by different patients’ and physicians’ attitudes. Although variation is not something negative by definition and is ‘natural’ up to a certain level, a better understanding of this phenomenon, addressing similarities and differences across specific settings and providers, should be pursued. Furthermore, in order to make informative comparisons among settings, there is need to standardize the measures used to measure the quantity and quality of antibiotic use.

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Transparency declarations

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Supplementary data
Tables S1–S9 are available as Supplementary data at JAC online.

References
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