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Escalating incidence of infective endocarditis in Europe in the 21st century

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ABSTRACT

Aim To provide a contemporary analysis of incidence trends of infective endocarditis (IE) with its changing epidemiology over the past two decades in Europe.

Methods A systematic review was conducted at the Mayo Clinic, Rochester. Ovid EBM Reviews, Ovid Embase, Ovid Medline, Scopus and Web of Science were searched for studies published between 1 January 2000 and 30 November 2020. All studies were independently reviewed by four referees and those that included a population-based incidence of IE in patients, irrespective of age, in Europe were included. Least squares regression was used to estimate pooled temporal trends in IE incidence.

Results Of 9138 articles screened, 18 studies were included in the review. Elderly men predominated in all studies. IE incidence increased 4.1% per year (95% CI 1.8% to 6.4%) in the pooled regression analysis of eight studies that included comprehensive and consistent trends data. When trends data were weighted according to population size of individual countries, an increase in yearly incidence of 0.27 cases per 100 000 people was observed. Staphylococci and streptococci were the most common pathogens identified. The rate of surgical intervention ranged from 10.2% to 60.0%, and the rate of inpatient mortality ranged from 14.3% to 17.5%. In six studies that examined the rate of injection drug use, five of them reported a rate of less than 10%.

Conclusion Based on findings from our systematic review, IE incidence in Europe has doubled over the past two decades in Europe. Multiple factors are likely responsible for this striking increase.

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Key questions

What is already known about this subject?

► Infective endocarditis (IE) is an uncommon life-threatening infection that, despite aggressive medical and surgical interventions, remains associated with high morbidity and mortality. Recent studies, however, have reported mixed results regarding temporal changes in trends of IE incidence.

What does this study add?

► This study pooled data from all nationwide population-based registries in Europe and demonstrated an alarming increase in IE incidence over the past two decades.

How might this impact on clinical practice?

► Our findings alert clinicians as to the increasing incidence of IE and the need to include it in the differential diagnosis of patients who present with systemic complaints of infection. The study also highlights the inefficiency in coding practices, which can compromise accurate IE incidence trends determinations.

INTRODUCTION

Infective endocarditis (IE) is one of the most lethal infection syndromes. Despite almost universal hospitalisation for initial treatment and the availability of multiple recent advances in diagnosis and management, it is characterised by a 1-year mortality rate that exceeds 30%.[1] Due to the rarity of the syndrome, both the management of, and investigation into the disease can be difficult. For the former, an IE diagnosis may be delayed, and treatment may be suboptimal resulting in worse outcomes. For the latter, generation of clinical trial data with adequate enrolment of patients can be challenging.

Fortunately, multiple European countries systematically record nationwide hospital admissions data and code the reasons for those admissions, which permit population-based investigations of the epidemiology of diseases.[2–4] Incidence trends, in particular, have been of keen interest and have been prompted, in part, by changes in the 2008 National Institute for Health and Care Excellence (NICE) guidelines[5] and the 2009 European Society of Cardiology (ESC) guidelines,[6] that either called for a total elimination of antibiotic prophylaxis (AP) use in the dental and other settings (NICE) or a marked reduction in this practice (ESC). As a result, concerns that an increase in IE
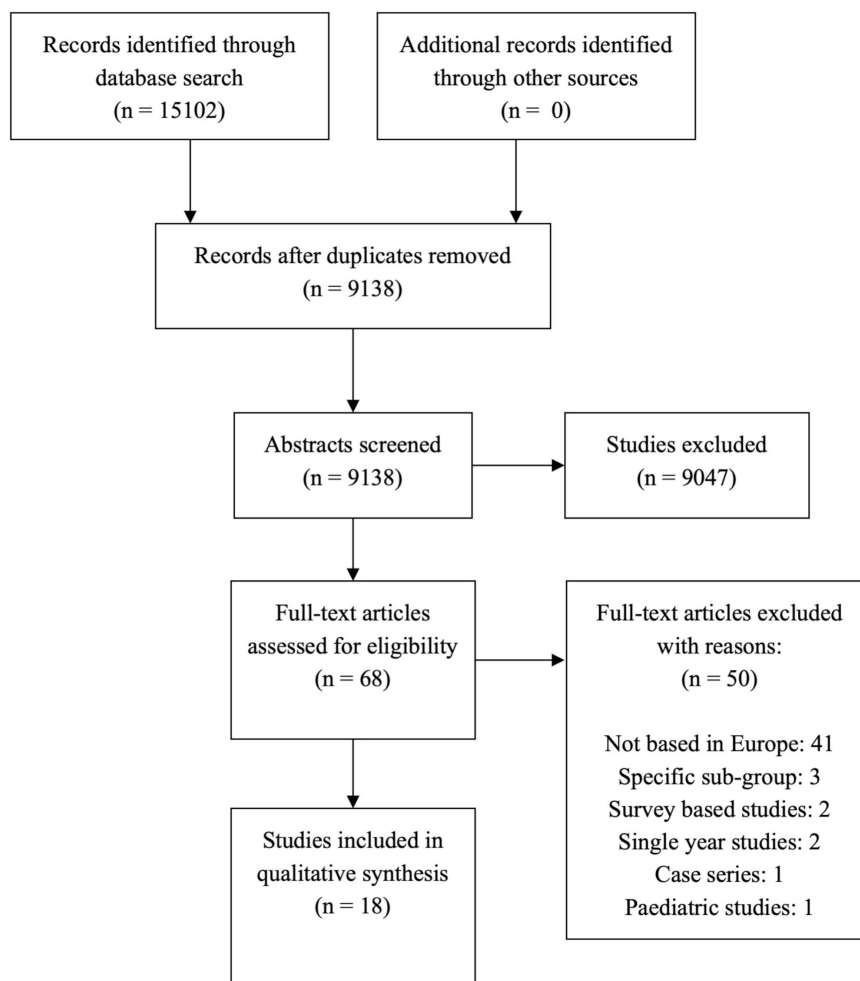
Figure 1 Schematic representation of study selection using PRISMA checklist. PRISMA, Preferred Reporting Items for Systematic Reviews and Meta-Analyses.

incidence due to viridans group streptococci (VGS) would occur following publication of these recommendations prompted investigations of nationwide data.

There are several other factors including ageing populations, increased comorbid conditions, increased placement of medical devices and injection drug use (IDU) that impact IE incidence trends and epidemiology. These factors have been addressed, to some degree, in most of the European national investigations.

Based on the availability of trend data of IE incidence and epidemiology from several countries in Europe, we conducted a contemporary systematic review of European population-based investigations of IE during the 21st century.

METHODS

A literature search was performed in December 2020 with a focus on the incidence and epidemiology of IE using Ovid EBM Reviews, Ovid Embase, Ovid Medline, Scopus and Web of Science to identify articles published between 1 January 2000 and 30 November 2020. The search was limited to the English language and search strategies are outlined in online supplemental material.

All results were exported to Endnote where duplicates were deleted. Two authors (KMT and LMB) performed the literature review and any disagreements were solved by discussion with two authors (MJD and DCD). Corresponding authors of studies were contacted via email in cases where queries existed.

Inclusion and exclusion criteria

All publications that provided information on population-based trends of IE in European populations from the year 2000 onwards were included in the review. Single-centre and multicentre studies, clinical trials, case reports, abstracts, systematic reviews and animal studies were excluded. Investigations that examined IE incidence specific to infecting pathogens or unique patient populations (eg, HIV, congenital heart disease) were excluded. Studies that included data from the COVID-19 pandemic period (January 2020 onwards) and those that reported data for less than 6 months each year for two or more years were excluded.

Data extraction

Data that described authors, publication year, study location, population covered, mean/median age, sex

Figure 2 Visual representation of European countries included in the systematic review.

distribution, incidence, microbiology, mortality, IDU and rates of surgical valvular intervention were extracted from all studies. Four authors (MD, KMT, VA and WT) worked independently to extract data from studies and contacted study investigators if additional data was required.

Study definition and outcomes
The primary outcome was the trend of IE incidence; secondary outcomes included temporal trends of pathogen prevalence, age, sex, prosthetic valve placement, IDU and mortality (in-patient, 30 days, 6 months and 1 year).

Risk of bias
Two reviewers (KMT and WT) independently rated the methodological quality of each study. The quality of each population-based survey was assessed, based on four key features: adequacy of population definition, sampling techniques, disease definition and completeness of case ascertainment (online supplemental table 1).[7] A population definition was deemed to be inadequate if the residency status of all IE patients was not ascertained. Since all studies used population data from national or regional registries, the population definition was deemed adequate for all studies. Optimal sampling techniques included complete enumeration or random sampling techniques. Disease definition was defined as adequate if studies used Duke/modified Duke criteria for a diagnosis of IE. Adequacy of case ascertainment to include all cases in a given country was assessed based on case-finding

procedures, inclusion of postmortem diagnoses and number of hospitals serving the population under study that participated in the study. Author statements about shortfall in case ascertainment were also considered an indication of inadequate case ascertainment. Based on these criteria, studies were excluded that had considerable shortfalls in case ascertainment and/or lacked a case definition. Reviewer disagreements were resolved by consensus after rereview of the article.

The study was registered with the International Prospective Register of Systematic Reviews, which is an international database of prospectively registered systematic reviews in health and social care (Registration ID: CRD42020191196).[8]

Statistical analysis
Studies included in the trend analysis had cases from more than two time points, at least 100 observed cases in each time period, and estimates of population size from which cases were observed. For each time point within a study, the incidence per 100 000 population was calculated by taking the observed cases divided by the population estimate multiplied by 100 000. Least squares regression was done on both the incidence per 100 000 and the log transformed incidence per 100 000. Interpretation of the regression results for the untransformed incidence yields an annual increase in cases per 100 000 while interpretation of the log transformed incidence per 100 000 resulted in an estimate of yearly percent increase

Table 1 Clinical features of patient populations in included studies

#	Author	Total cases (N)	Years included	Country	AGE	Male sex %	Microbiology (%)	Prosthetic valve %	Cardiovascular device %	INJECTION DRUG USE %	Mortality %	Required surgery %
1	Scudeller 2009[18]	254	2004–2008	Italy	Mean: 67.0 (SD 14.0)	66.5	Enterococci 21.6 *S. aureus* 17.5 VGS 14.0	32.3	NP	2.0	3 months 20.5	40.2
2	Fedeli 2011[26]	1863	2000–2008	Italy	Median 68.0 (IQR: 57.0–77.0)	63.0	Blood culture data *S. aureus* 29.5 CoNS 6.7 *Enterococcus faecalis* 12.4 Streptococci 28.6 Secondary code data Streptococci 36.7 *S. aureus* 17.4 CoNS 6.0 *E. faecalis* 5.3	6.2	NP	NP	Overall 14.3	In-patient 37.0 1 year 38.0
3	Thornhill 2011[14]		2000–2010	England	NP	NP	NP	NP	NP	NP	NP	NP
4	Ternhag 2013[16]	7609	1997–2007	Sweden	Mean 65.7 (IQR: 55–79)	59.2	NP	11.7	NP	4.7	30 days—33.7 5 years—14.7	5 years follow-up 13.0
5	Dayer 2015[2]	19804	2000–2013	England	Mean 59.1 59.0 (SD: 20.3) before 2008, 59.3 (SD 20.8) after 2008	68.5	NP	NP	NP	NP	NP	NP
6	Cresti 2016[1]	170	1998–2014	Italy	Mean 65.7(SD: 16.0)	60.9	S. *aureus* 25.0 CoNS 22.0 *Streptococcus viridans* 15.0 Enterococci 14.0	30.0	NP	4.0	1 year 31.8 In-patient 24.0	1 year 46.5; Urgent/emergent 31.8; Operated within 10 days 29.0
7	Erichsen 2016[24]	5486	1994–2011	Denmark	Mean: 62.7 (no SD provided)	64.4	NP	NP	NP	NP	NP	NP
8	Keller 2016[3]	94364	2005–2014	Germany	NP	NP	Streptococci 20.8 Staphylococci 21.9	NP	NP	NP	Overall 17.0	NP
9	van den Brink 2016[31]	5213	2005–2011	The Netherlands	Mean 67.5 (range: 22.0–97.0)	69.9.	Staphylococci 36.1 *S. aureus* 30.1 Streptococci 37.4	30.1	7.9	NP	All-cause 36.1% (Median follow-up 4.2 years)	38.9
10	Olmos 2017[17]	16867	2003–2014	Spain	Mean 63.8 (SD: 17.5)	66.3	Streptococci 20.4 *S. aureus* 17.1 Enterococci 13.1 CoNS 12.2	18.0	1.1	2.6	In-patient 20.4	23.0
11	Ahtela 2018[11]	2611	2005–2014	Finland	Mean 60.0 (SD: 18.3)	68.2	NP	5.5	1.5	NP	30 days 11.3	NP
12	Jordal 2018[12]	706	1996–2015	Norway	Mean 59.2 (need to calculate combined SD)	69.1	*S. aureus* 31.1 VGS 23.1 CoNS 11.5 Enterococci 9.5	29.9	NP	20.7	30 days 12.7 1 year 21.4	34.1
13	Ortega 2019[23]	25952	1997–2014	Spain	Overall mean: 62.2 (18.6)	65.9	Staphylococci 32.7 *S. aureus* 19.6 Streptococci 2.5	9.6	6.5; CIED related IE 21.2	6.8	90 days 26.2	NP
14	Jensen 2020[4]	7669	1997–2017	Denmark	Median 70.2 (IQR: 58.3–78.8)	65.2	NP	15.9	10.9	NP	NP	NP

Continued

Table 1 Continued

#	Author	Total cases (N)	Years included	Country	AGE	Male sex %	Microbiology (%)	Prosthetic valve %	Cardiovascular device %	INJECTION DRUG USE %	Mortality %	Required surgery %
15	Quan 2020[25]		1998–2017	England	NP	NP	Streptococci 49.0 Staphylococci 43.0	NP	NP	NP	NP	NP
16	Shah 2020[13]	7638	1990–2014	Scotland	Mean 65.0 (SD: 17.0)	49.0	Staphylococci 42.4 S. aureus 31.7 Streptococci 35.5 Enterococci 8.9	NP	NP	NP	1 year 32.0 30 days 14.7%	30 days 4.8 1 year 10.6
17	Thornhill 2020[15]		1998–2018	England	NP	NP	NP	NP	NP	NP	NP	NP
18	Vähäsarja[10]	4647	2008–2017	Sweden	Mean 64.9 (range: 17.0–100.0)	68.0	S. aureus 37.0 VGS 24.7%	NP	NP	NP	NP	In-patient 10.2

*The genus and species of the pathogens have been listed as presented in the individual studies. Since the pathogens were grouped differently in each study, it was not possible for us to standardise them.
CIED, cardiac implantable electronic device; CoNS, coagulase-negative staphylococci; IE, infective endocarditis; NP, not provided; VGS, viridans group streptococci.

in incidence rates. SEs and 95% confidence limits were calculated.

The pooled regression estimate from the primary analysis to the simple unweighted average of the individual regression estimates were compared and were similar. This suggested that the pooled regression was a reasonable estimate of the overall trend.

Patient and public involvement
No patients or public were involved at any stage of the synthesis of this study.

RESULTS
Study selection
A total of 9138 studies were identified from the search engines after deduplication. The study abstracts were screened, and 91 were identified for full text review. Eighteen studies met our inclusion criteria. A schematic representation of studies included using Preferred Reporting Items for Systematic Reviews and Meta-Analyses guidelines[9] is included in figure 1. England had the highest number of studies included (4); the remainder of geographical distribution of included studies is illustrated in figure 2. A detailed profile of studies is presented in table 1. A total of 15 studies defined IE using a primary or secondary diagnosis that was based on International Classification of Diseases (ICD), ninth revision (ICD 9) and tenth revision (ICD 10) (table 2).

Patient demographics
IE was predominately seen in older patients with the lowest mean age (59.1 years) recorded by Dayer[2] and the highest median age (70.2) recorded by Jensen.[4] Fourteen studies included investigations included all patient age groups. Vähäsarja et al[10] included patients aged ≥17 years, Ahtela et al[11] and Jordal et al[12] included patients aged ≥18 years and Shah[13] included patients aged ≥20 years. IE predominantly affected men, as reported in all but one study.[13]

Overall incidence
All studies reported temporal trends in IE incidence as illustrated in figure 3. Results of three studies from England[2 14 15] were combined into one line graph since there was duplication of data owing to similar methodology and overlapping time periods. Overall, an appreciable increase in the incidence of IE was demonstrated.

Figure 4 displays estimates from each country with the pooled regression slope overlaid on individually observed data. Table 3 shows the least squares regression slopes for each study included in pooled log scale regression analysis for temporal trends of IE incidence and the average of individual studies. The pooled regression estimate was 4.1%±1.2% per year increase in IE incidence, amounting to a compound increase in incidence of 106% over 18 years. The two Swedish studies[10 16] used different databases and differed in methodology; hence, they were excluded from the primary analysis. They were

Table 2 List of ICD codes used in included studies

Study	ICD codes	Primary/secondary/tertiary
Scudeller 2009[18]	No ICD code data available in study	–
Fedeli 2011[26]	ICD 9: 421, 98.84 or 112.81	P+S
Thornhill 2011[14]	ICD 10: I33	P+S
Ternhag 2013[16]	ICD 10: I33, I38 or I39	Unspecified
Dayer 2015[2]	ICD 10: I33	P+S
Cresti 2016[1]	ICD 9: 421 .x	P+S
Erichsen 2016[24]	ICD 10: I33 or I38.9	Unspecified (first-time code of incident IE recorded)
Keller 2016[3]	ICD 10: I33	Unspecified
van den Brink 2016[31]	Insurance-based codes used for case extraction	–
Olmos 2017[17]	ICD 9: 421.0, 421.1, 421.9 or 424.99	P+S
Ahtela 2018[11]	ICD 10: I33, I38 or I39 (specificity 96.8%)	P (66.3%)+S (24.1%)+T (9.6%)
Jordal 2018[12]	ICD 10: I33.0, I38, or I39.0 (after year 1999)	Unspecified
Ortega 2019[28]	ICD 9: 421.0, 421.1, 421.9, 112.81, 115.04, 115.14, or 115.94	P+S
Jensen 2020[4]	ICD 10: I33.x, I38.x or I39.8 ICD 8: 421	P+S
Quan 2020[25]	ICD 10: I33.0, I33.9, I39.0, I39.8, I01.1, B37.6, or T82.6	P+S
Shah 2020[13]	ICD 10: I33, I38 or I39 ICD 9: 421.1, 424.91, 424.90 or 424.99	Unspecified
Thornhill 2020[15]	ICD 10: I33	P+S
Vähäsarja 2020[10]	No ICD code data available in study	–

ICD, International Classification of Diseases.

included in online supplemental table 2 and figure 1, which demonstrated an overall estimated percent annual increase of 2.8%±1.0%. The Norway study by Jordal et al. was also excluded as it only provided two data points of decade-wise estimates of IE cases and did not include yearly population estimates used to calculate annual incidence.

Finally, we investigated the degree to which the pooled trend might be altered if we weighted regression estimates based on the underlying country populations. This was done on original scale, incidence per 100 000. Because all studies had estimates from 2008, the population in 2008 was used as the weight in this calculation. As shown in table 3, the estimated yearly increase in IE incidence per 100 000 was 0.24 IE cases per 100 000 per annum without weighting. When weights were applied this increased to 0.27 IE cases per 100 000 per annum. This is likely due to the German study[3] having the highest estimated yearly
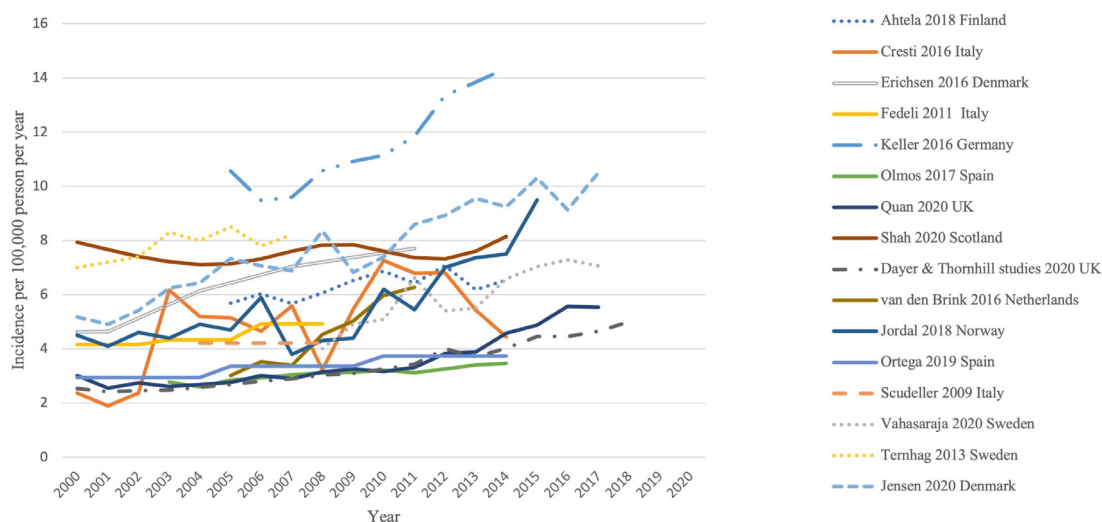

Figure 3 Temporal trends of crude incidence of IE across all studies from 2000 to 2020. The y-axis denotes number of cases per 100 000 people while the x-axis denotes years 2000–2020. IE, infective endocarditis.
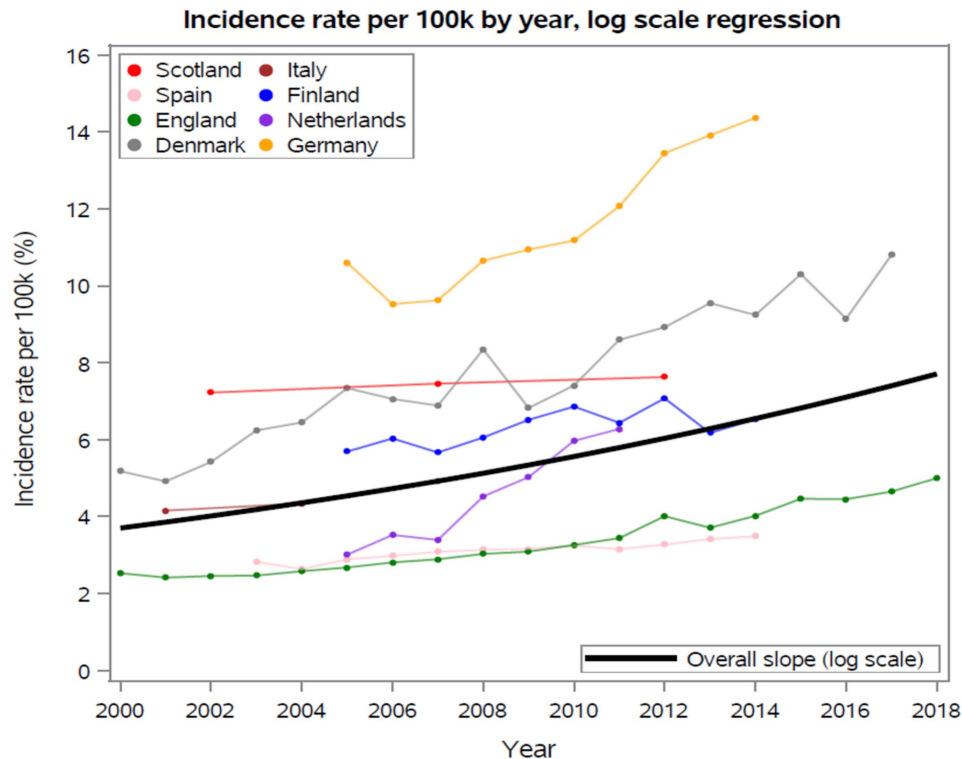
Figure 4 Individual and pooled incidence rate per 100 000/year, log scale regression. The y-axis denotes incidence rate per 100 000 (%), while the x-axis denotes years 2000–2018.

increase as well as the highest population. If Germany data were excluded, the unweighted average was 0.20 IE cases per 100 000 per annum and the weighted average was 0.16 IE cases per 100 000 per annum.

Injection drug use
Six studies reported percentages of patients with IDU and IE (table 1). Trends in IDU over study periods were reported by two other studies. Jordal et al[12] reported an overall increase in IDU from 16.5% to 23.5% from 1996

to 2015. In contrast, Olmos et al[17] reported an overall decrease from 4.4% to 1.1% from 2003 to 2014 in Spain.

Microbiology
Details of microbiological data were provided in 11 studies (table 1). Seven studies reported staphylococcal species as the most common pathogens, of which three specified *Staphylococcus aureus* as the most common organism. Streptococcal species were identified as the most common pathogens in three studies. Scudeller et

Table 3 Least squares regression slopes for each study included in the pooled log scale regression analysis for temporal trends of IE incidence and the average of individual studies

| Studies included | Country | Incidence per 100 k | | Log incidence | |
		Estimated annual increase (per 100 k)	95% CI	Estimated annual increase (%)	95% CI
Shah 2020[13]	Scotland	0.040	0.033 to 0.047	0.54	0.44 to 0.64
Olmos 2017[17]	Spain	0.064	0.050 to 0.078	2.09	1.61 to 2.57
Dayer and Thornhill 2011–2020[2]	England	0.144	0.126 to 0.162	4.23	3.82 to 4.64
Jensen 2020[4]	Denmark	0.312	0.263 to 0.361	4.17	3.50 to 4.84
Fedeli 2011[26]	Italy	0.128	0.052 to 0.204	2.83	1.58 to 4.08
Ahtela 2019[11]	Finland	0.103	0.024 to 0.182	1.66	0.42 to 2.89
van den Brink 2016[31]	Netherlands	0.583	0.470 to 0.696	13.05	10.59 to 15.51
Keller 2016[3]	Germany	0.535	0.387 to 0.683	4.52	1.77 to 6.39
	Average of individual studies	0.239	0.060 to 0.418	4.14	0.90 to 7.37
	Pooled regression	0.238	0.096 to 0.380	4.08	1.77 to 6.39

IE, infective endocarditis.

al[18] reported enterococci as the most common causative pathogens. Olmos et al[17] demonstrated a significant rise in yearly percentage prevalence of enterococci from 10.4% in 2003 to 16.6% in 2014.

Outcomes

Details of the proportion who underwent surgery are listed in table 1 (range, 10.2%– 60.0%). Follow-up duration among these patients was highly variable (inpatient to 5 year post-IE diagnosis). Other outcomes examined included inpatient, 30 days, 3 months and 1-year mortality rates (table 1).

DISCUSSION

This is the first systematic review that has included pooled trends of incidence of IE across multiple nationwide population-based studies in Europe. The current investigation has highlighted a 4% per year rise in incidence of IE which, when compounded, is an alarming doubling in incidence between 2000 and 2018. Based on findings of our extensive systematic review that involved multiple countries with nationwide databases, support the notion that the IE incidence escalation seen is valid and is likely due to multiple factors operative in the 21st century. Factors that need to be considered include (1) improvements in diagnosis; (2) changes in epidemiology and associated risk factors; (3) restrictions in AP use promulgated by updated versions of guidelines and (4) improvements in coding practice.

Diagnostic advances in IE have characterised the past two decades and, while not examined specifically in our review, these advances likely contributed to the increasing incidence of IE. Li et al[19] modified the Duke criteria that included echocardiography as a pivotal tool in establishing an IE diagnosis in patients who do not undergo valve surgery or autopsy for diagnosis confirmation. Inpatient data available through the Nationwide Inpatient Sample for the first decade of this century in the USA supports the notion that echocardiography use has indeed increased among hospitalised patients[20 21] and anecdotally the use of echocardiography is far more widespread in Europe too. Echocardiography, both transthoracic and transoesophageal has become a 'mainstay' of cardiovascular evaluation in suspect cases of IE and its complications. The increase in use of echocardiography, particularly the transoesophageal approach is expected to continue which is a potential factor in the increase in IE incidence due to enhanced diagnostic features.

Additional tools, in particular multislice CT, [18]fluorodeoxyglucose positron emission tomography/CT, and single-photon emission CT, have also been helpful in securing an IE diagnosis when transoesophageal echocardiography has been insufficient or unavailable. Such methods were incorporated in the latest ESC guidelines.[6] In addition, there have been significant laboratory advances in performing blood cultures, which are also key in the diagnosis of IE using the modified Duke criteria.[22]

Studies included in this review did not provide an analysis of factors that could have accounted for the increase in IE incidence and a temporal sequence of prevalence of epidemiological variables. It is, however, intuitive that contemporary aspects of healthcare that include invasive procedures and placement of indwelling cardiovascular devices have increased over the past two decades and likely impacted both IE incidence and its clinical features. Nevertheless, four investigations documented an increase in prosthetic valve placement.[4 17 23 24]

Although some studies have reported increasing IE incidence after guideline changes,[23] this was not found universally. It would be premature to link a rise in IE incidence in Europe to restriction of AP use. Quan et al reported, for example, increasing trends of IE in England which neither correlated with the change in recommendations by NICE in 2008, nor to a rise in prevalence of VGS as a causative pathogen.[25] Moreover, most of the included studies in this systematic review identified staphylococci and streptococci as the most common pathogens of IE over the past two decades (table 1). Hence, an increase in IE incidence cannot be explained simply by trends in pathogen prevalence, which has its own array of pitfalls. This includes the fact that not all patient records included secondary designated microbiology codes, which provides a partial profile of microbiology. Fedeli et al noted a shockingly low percentage (~21%) of secondary code designations of IE-related pathogens in Italy.[26] Dayer et al reported that ~50% of cases in England were designated secondary codes for microbiology, which increased towards the end of the study period in 2013.[2] Similarly, Quan et al reported that only 67% of the total cases were designated codes for microbiology.[25] Perhaps the most important inherent limitation in all studies was the absence of a specific ICD-10 code for VGS. This has greatly hampered accuracy of reporting microbiological data as we strive to define the impact, if any, of changes in AP use in the prevention of IE.

The impact of ICD coding on IE incidence is often underappreciated and it is critical to emphasise because many of the investigations included in our review were dependent on ICD coding to identify IE admissions (table 2). Cresti et al, for example, found that 28% of IE cases extracted using ICD-9 codes were false positive and failed to pick up 14% of confirmed cases, which raised questions on the validity of current code designations.[1] The potential impact of coding was also highlighted by Fawcett et al[27] who reported a sensitivity of IE of 76% for specific codes in ICD-10 with more than half of cases coded by using ICD-10 as IE were not confirmed cases. The code I33 had a positive predictive value (PPV) of 82%–85%; in contrast, and the code I38 had a PPV of <6% and accounted for many of the false-positive cases and was used in seven investigations in the current systematic review. Although such coding practices may not have a major impact on estimating burden of disease for more

prevalent diseases, for IE, an uncommon malady, consistent coding practices to accurately estimate temporal trends is essential. Because 12 of 23 studies that were included in our systematic review used ICD-10 coding and all included I33, the most specific code for IE, significant trend variation in IE incidence across investigations was not observed (figure 2).

The two Spanish investigations[17 23] included in the systematic review reported incidence data for a similar time period. Ortega-Loubon *et al* included a larger sample size and a higher annual incidence but reported a similar trend to the Olmos study. However, Ortega-Loubon *et al* reported a higher prevalence (16%) of Gram-negative rods as compared with streptococcal species (2.5%), while Olmos *et al* identified streptococcal species as the most common causative pathogen of IE (20.4%). Two similar contemporary studies with the same methodology from the authors of the Ortega study reported that causative pathogens were 'unspecified' in ~86% of cases,[28 29] while the 2019 study reported microbiological findings in 52.2% of cases. This suggests that microbiological data from the Ortega study may not be reliable and Olmos *et al* likely provided a more accurate representation of the pathogen distribution in Spain.

Williams *et al* recently published a systematic review detailing contemporary epidemiological changes in IE following major guideline changes.[30] They included studies from North America and Europe and concluded that although there was no appreciable increase in IE incidence in North America following pertinent guideline changes, there was a potential rise in incidence in Europe. They included just five European studies in their review, whereas our current investigation includes 18 studies, where the incidence trends were examined statistically over 18 years. Furthermore, Williams *et al* specifically studied the impact of guideline changes for AP use on the incidence of IE, while we focused on trends, irrespective of AP guideline changes, to assess all factors responsible for increasing IE trends.

Despite the thoroughness of this review, there were some limitations to the current study. Data for patient demographics, microbiology, mortality and surgery were provided for most studies as a single percentage over the study period, and not as yearly trends. Hence, a risk factor meta-regression analysis could not be performed, which would have helped quantify the contribution of specific risk factors associated with the rising trends of IE. Furthermore, these studies are only observational and by design, unable to determine aetiological factors associated increasing incidence.

CONCLUSION

This study highlights a greater than twofold increase in incidence of IE in Europe between 2000 and 2018. This is likely due to a combination of factors including an increasingly elderly population, a rise in cardiovascular device implantation procedures, increased use of multiple imaging modalities for diagnosis, improved coding and possibly a restriction of AP use. However, it is difficult to ascertain a single factor for this change, since there is great heterogeneity in how data are reported across studies and inadequacy in coding of microbiological data.

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REFERENCES
1 Cresti A, Chiavarelli M, Scalese M, *et al*. Epidemiological and mortality trends in infective endocarditis, a 17-year population-based prospective study. *Cardiovasc Diagn Ther* 2017;7:27–35.
2 Dayer MJ, Jones S, Prendergast B, *et al*. Incidence of infective endocarditis in England, 2000-13: a secular trend, interrupted time-series analysis. *Lancet* 2015;385:1219–28.

3 Keller K, von Bardeleben RS, Ostad MA, *et al*. Temporal trends in the prevalence of infective endocarditis in Germany between 2005 and 2014. *Am J Cardiol* 2017;119:317–22.

4 Jensen AD, Bundgaard H, Butt JH, *et al*. Temporal changes in the incidence of infective endocarditis in Denmark 1997-2017: a nationwide study. *Int J Cardiol* 2021;326:145–52.

5 Richey R, Wray D, Stokes T, *et al*. Prophylaxis against infective endocarditis: summary of NICE guidance. *BMJ* 2008;336:770–1.

6 Habib G, Lancellotti P, Antunes MJ, *et al*. 2015 ESC guidelines for the management of infective endocarditis: the task force for the management of infective endocarditis of the European Society of Cardiology (ESC). endorsed by: European Association for Cardio-Thoracic Surgery (EACTS), the European Association of Nuclear Medicine (EANM). *Eur Heart J* 2015;36:3075–128.

7 Tleyjeh IM, Abdel-Latif A, Rahbi H, *et al*. A systematic review of population-based studies of infective endocarditis. *Chest* 2007;132:1025–35.

8 Page MJ, Shamseer L, Tricco AC. Registration of systematic reviews in PROSPERO: 30,000 records and counting. *Syst Rev* 2018;7:018–699.

9 Page MJ, McKenzie JE, Bossuyt PM. The PRISMA 2020 statement: an updated guideline for reporting systematic reviews. *Bmj* 2021;29.

10 Vähäsarja N, Lund B, Ternhag A, *et al*. Incidence of infective endocarditis caused by viridans group streptococci in Sweden - effect of cessation of antibiotic prophylaxis in dentistry for risk individuals. *J Oral Microbiol* 2020;12:1768342.

11 Ahtela E, Oksi J, Porela P, *et al*. Trends in occurrence and 30-day mortality of infective endocarditis in adults: population-based registry study in Finland. *BMJ Open* 2019;9:e026811.

12 Jordal S, Kittang BR, Salminen P-R, *et al*. Infective endocarditis in Western Norway: a 20-year retrospective survey. *Infect Dis* 2018;50:757–63.

13 Shah ASV, McAllister DA, Gallacher P, *et al*. Incidence, microbiology, and outcomes in patients hospitalized with infective endocarditis. *Circulation* 2020;141:2067–77.

14 Thornhill MH, Dayer MJ, Forde JM, *et al*. Impact of the NICE guideline recommending cessation of antibiotic prophylaxis for prevention of infective endocarditis: before and after study. *BMJ* 2011;342:d2392.

15 Thornhill MH, Dayer MJ, Nicholl J, *et al*. An alarming rise in incidence of infective endocarditis in England since 2009: why? *Lancet* 2020;395:1325–7.

16 Ternhag A, Cederström A, Törner A, *et al*. A nationwide cohort study of mortality risk and long-term prognosis in infective endocarditis in Sweden. *PLoS One* 2013;8:e67519.

17 Olmos C, Vilacosta I, Fernández-Pérez C, *et al*. The Evolving Nature of Infective Endocarditis in Spain: A Population-Based Study (2003 to 2014). *J Am Coll Cardiol* 2017;70:2795–804.

18 Scudeller L, Badano L, Crapis M, *et al*. Population-based surveillance of infectious endocarditis in an Italian region. *Arch Intern Med* 2009;169:1718–3.

19 Li JS, Sexton DJ, Mick N, *et al*. Proposed modifications to the Duke criteria for the diagnosis of infective endocarditis. *Clin Infect Dis* 2000;30:633–8.

20 Papolos A, Narula J, Bavishi C, *et al*. U.S hospital use of echocardiography: insights from the nationwide inpatient sample. *J Am Coll Cardiol* 2016;67:502–11.

21 Virnig BA, Shippee ND, O'Donnell B. *Trends in the use of echocardiography, 2007 to 2011: Data Points #20.*

22 Liesman RM, Pritt BS, Maleszewski JJ, *et al*. Laboratory diagnosis of infective endocarditis. *J Clin Microbiol* 2017;55:2599–608.

23 Ortega-Loubon C, Muñoz-Moreno MF, Andrés-García I, *et al*. Nosocomial vs. community-acquired infective endocarditis in Spain: location, trends, clinical presentation, etiology, and survival in the 21st century. *J Clin Med* 2019;8. doi:10.3390/jcm8101755. [Epub ahead of print: 22 10 2019].

24 Erichsen P, Gislason GH, Bruun NE. The increasing incidence of infective endocarditis in Denmark, 1994-2011. *Eur J Intern Med* 2016;35:95–9.

25 Quan TP, Muller-Pebody B, Fawcett N, *et al*. Investigation of the impact of the NICE guidelines regarding antibiotic prophylaxis during invasive dental procedures on the incidence of infective endocarditis in England: an electronic health records study. *BMC Med* 2020;18:020–1531.

26 Fedeli U, Schievano E, Buonfrate D, *et al*. Increasing incidence and mortality of infective endocarditis: a population-based study through a record-linkage system. *BMC Infect Dis* 2011;11:1471–2334.

27 Fawcett N, Young B, Peto L, *et al*. 'Caveat emptor': the cautionary tale of endocarditis and the potential pitfalls of clinical coding data-an electronic health records study. *BMC Med* 2019;17:019–1390.

28 Ortega-Loubon C. Ruiz López del Prado G, Muñoz-Moreno MF, et al. impact of the economic crisis on endocarditis mortality in Spain: a nationwide study. *Int J Health Serv* 2021;29:00207314211012357.

29 Heredia-Rodríguez M, Hernández A, Bustamante-Munguira J, *et al*. Evolution of the incidence, mortality, and cost of infective endocarditis in Spain between 1997 and 2014. *J Gen Intern Med* 2018;33:1610–3.

30 Williams ML, Doyle MP, McNamara N, *et al*. Epidemiology of infective endocarditis before *versus* after change of international guidelines: a systematic review. *Ther Adv Cardiovasc Dis* 2021;15:17539447211002687.

31 van den Brink FS, Swaans MJ, Hoogendijk MG, *et al*. Increased incidence of infective endocarditis after the 2009 European Society of Cardiology guideline update: a nationwide study in the Netherlands. *Eur Heart J Qual Care Clin Outcomes* 2017;3:141–7.

APPENDIX

ONLINE SEARCH STRATEGY:

EBM Reviews:
((incidence* or epidemio*).ab,fs,hw,ti or (((number or new) adj2 cases) or surveillance or biosurveillance or pharmacoepidemiolog* or seroepidemio* or (morbidity adj2 (rate* or pattern*)) or infection-rate* or sampl* or age-distribution or sex-distribution or geographic-distribution* or case-mix or community-assessment* or disease-registr* or prevalence).ab,fs,hw,ti.) AND (endocarditis.hw,ti. or (endocarditis adj5 (incidence* or epidemio* or cases or surveillance or biosurveillance or pharmacoepidemiolog* or seroepidemio* or rate* or prevalence)).ab,hw,ti.) NOT ("case report*".pt,ti.) Limit to English, 2000+

Embase (1974+)
(exp incidence/ or exp epidemiological data/ or exp epidemiology/ or (((number or new) adj2 cases) or surveillance or biosurveillance or pharmacoepidemiolog* or seroepidemio* or (morbidity adj2 (rate* or pattern*)) or infection-rate* or sampl* or age-distribution or sex-distribution or geographic-distribution* or case-mix or community-assessment* or disease-registr* or prevalence).ab,kw,ti.) AND (endocarditis/ or bacterial endocarditis/ or fungal endocarditis/ or prosthetic valve endocarditis/ or subacute endocarditis/ or endocarditis.kw,ti. or (endocarditis adj5 (incidence* or epidemio* or cases or surveillance or biosurveillance or pharmacoepidemiolog* or seroepidemio* or rate* or prevalence)).ab.) NOT (exp animal/ not exp human/, 12 "case report".kw,pt,ti.) Limit to English, 2000+, article or article in press

Ovid MEDLINE(R) 1946 to Present and Epub Ahead of Print, In-Process & Other Non-Indexed Citations and Ovid MEDLINE(R) Daily:
(Incidence/ or exp Epidemiology/ or exp Population Surveillance/ or (incidence* or epidemio*).ab,hw,kf,ti,fx. or (((number or new) adj2 cases) or surveillance or biosurveillance or pharmacoepidemiolog* or seroepidemio* or (morbidity adj2 (rate* or pattern*)) or infection-rate* or sampl* or age-distribution or sex-distribution or geographic-distribution* or case-mix or community-assessment* or disease-registr* or prevalence).ab,kf,ti.) AND (Endocarditis, Bacterial/ or Endocarditis/ or Endocarditis, Subacute Bacterial/ or endocarditis.kw,ti. or (endocarditis adj5 (incidence* or epidemio* or cases or surveillance or biosurveillance or pharmacoepidemiolog* or seroepidemio* or rate* or prevalence)).ab.) NOT (exp Animals/ not Humans/, "case report".kw,pt,ti.) Limit to English, 2000+

Scopus:
(((TITLE (endocarditis) OR KEY (endocarditis) AND TITLE-ABS-KEY (endocarditis W/5 (incidence* OR epidemio* OR cases OR surveillance OR biosurveillance OR pharmacoepidemiolog* OR seroepidemio* OR rate* OR prevalence)))) AND ((TITLE (endocarditis) OR KEY (endocarditis) AND TITLE-ABS-KEY (endocarditis W/5 (incidence*

OR epidemio* OR cases OR surveillance OR biosurveillance OR pharmacoepidemiolog* OR seroepidemio* OR rate* OR prevalence))))) AND NOT (TITLE ("case report*" OR animal* OR rat OR rats OR mice OR mouse OR rabbit* OR rodent*)) AND (LIMIT-TO (DOCTYPE , "ar") OR LIMIT-TO (DOCTYPE , "re") OR LIMIT-TO (DOCTYPE , "er")) AND (LIMIT-TO (LANGUAGE , "English")) AND (LIMIT-TO (SRCTYPE , "j")) AND (LIMIT-TO (PUBYEAR , 2020) OR LIMIT-TO (PUBYEAR , 2019) OR LIMIT-TO (PUBYEAR , 2018) OR LIMIT-TO (PUBYEAR , 2017) OR LIMIT-TO (PUBYEAR , 2016) OR LIMIT-TO (PUBYEAR , 2015) OR LIMIT-TO (PUBYEAR , 2014) OR LIMIT-TO (PUBYEAR , 2013) OR LIMIT-TO (PUBYEAR , 2012) OR LIMIT-TO (PUBYEAR , 2011) OR LIMIT-TO (PUBYEAR , 2010) OR LIMIT-TO (PUBYEAR , 2009) OR LIMIT-TO (PUBYEAR , 2008) OR LIMIT-TO (PUBYEAR , 2007) OR LIMIT-TO (PUBYEAR , 2006) OR LIMIT-TO (PUBYEAR , 2005) OR LIMIT-TO (PUBYEAR , 2004) OR LIMIT-TO (PUBYEAR , 2003) OR LIMIT-TO (PUBYEAR , 2002) OR LIMIT-TO (PUBYEAR , 2001) OR LIMIT-TO (PUBYEAR , 2000))

Web of Science:
(TOPIC: (incidence* or epidemio*) *OR* TOPIC: ((((number or new) NEAR/2 cases) or surveillance or biosurveillance or pharmacoepidemiolog* or seroepidemio* or (morbidity NEAR/2 (rate* or pattern*)) or infection-rate* or sampl* or age-distribution or sex-distribution or geographic-distribution* or case-mix or community-assessment* or disease-registr* or prevalence)) AND TOPIC: (endocarditis) *OR* TOPIC: (endocarditis NEAR/5 (incidence* or epidemio* or cases or surveillance or biosurveillance or pharmacoepidemiolog* or seroepidemio* or rate* or prevalence))) NOT TI=("case report*" OR animal* OR rat OR rats OR mice OR mouse OR rabbit* OR rodent*), Limit to English, Article, 2000+

Table 1: Quality assessment of included studies.

Study	Adequacy of population definition	Sampling techniques	Disease definition	Completeness of case ascertainment
Scudeller 2009	Adequate	Adequate	Adequate	Adequate
Fedeli 2011	Adequate	Adequate	Inadequate	Adequate
Thornhill 2011	Adequate	Adequate	Inadequate	Adequate
Ternhag 2013	Adequate	Adequate	Inadequate	Adequate
Dayer 2015	Adequate	Adequate	Inadequate	Adequate
Cresti 2016	Adequate	Adequate	Adequate	Adequate
Erichsen 2016	Adequate	Adequate	Inadequate	Adequate
Keller 2016	Adequate	Adequate	Inadequate	Adequate
van den Brink 2016	Adequate	Adequate	Inadequate	Adequate
Olmos 2017	Adequate	Adequate	Inadequate	Adequate
Ahtela 2018	Adequate	Adequate	Adequate	Adequate
Jordal 2018	Adequate	Adequate	Adequate	Adequate
Ortega 2019	Adequate	Adequate	Inadequate	Adequate
Jensen 2020	Adequate	Adequate	Inadequate	Adequate
Quan 2020	Adequate	Adequate	Inadequate	Adequate
Shah 2020	Adequate	Adequate	Inadequate	Adequate
Thornhill 2020	Adequate	Adequate	Inadequate	Adequate
Vähäsarja 2020	Adequate	Adequate	Adequate	Adequate

Table 2. Least squares regression slopes for each study in pooled log scale regression analysis for temporal trends of IE incidence, including Swedish studies.

Studies Included	Country	Incidence per 100k		Log incidence	
		Estimated annual increase (per 100k)	95% CI*	Estimated annual increase (%)	95% CI*
Shah 2020	Scotland	0.040	0.033-0.047	0.54%	0.44-0.64%
Olmos 2017	Spain	0.064	0.050-0.078	2.09%	1.61-2.57%
Dayer & Thornhill 2011-2020	England	0.144	0.126-0.162	4.23%	3.82-4.64%
Jensen 2020	Denmark	0.312	0.263-0.361	4.17%	3.50-4.84%
Fedeli 2011	Italy	0.128	0.052-0.204	2.83%	1.58-4.08%
Ahtela 2019	Finland	0.103	0.024-0.182	1.66%	0.42-2.89%
van den Brink 2016	Netherlands	0.583	0.470-0.696	13.05%	10-59-15.51%
Ternhag 2013	Sweden (2000-2007)	0.169	0.055-0.283	2.21%	0.74-3.66%
Vähäsarja 2020	Sweden (2008-2017)	0.327	0.209-0.445	5.76%	3.63-7.89%
Keller 2016	Germany	0.535	0.387-0.683	4.52%	1.77-6.39%
	Average of individual studies	0.241	0.103-0.378	4.11%	1.59-6.62%
	Pooled regression	0.161	0.043-0.279	2.81%	0.85-4.77%

*CI: Confidence Interval

Figure 1: Individual and pooled incidence rate per 100,000/year, log scale regression, including Swedish studies.