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ORIGINAL RESEARCH

Prevalence and incidence rates of atrial fibrillation in Norway 2004–2014

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ABSTRACT

Objective To study time trends in incidence of atrial fibrillation (AF) in the entire Norwegian population from 2004 to 2014, by age and sex, and to estimate the prevalence of AF at the end of the study period.

Methods A national cohort of patients with AF (≥ 18 years) was identified from inpatient admissions with AF and deaths with AF as underlying cause (1994–2014), and AF outpatient visits (2008–2014) in the Cardiovascular Disease in Norway (CVDNOR) project. AF admissions or out-of-hospital death from AF, with no AF admission the previous 10 years defined incident AF. Age-standardised incidence rates (IR) and incidence rate ratios (IRR) were calculated. All AF cases identified through inpatient admissions and outpatient visits and alive as of 31 December 2014 defined AF prevalence.

Results We identified 175 979 incident AF cases (30% primary diagnosis, 69% secondary diagnosis, 0.6% out-of-hospital deaths). AF IRs (95% confidence intervals) per 100 000 person years were stable from 2004 (433 (426–440)) to 2014 (440 (433–447)). IRs were stable or declining across strata of sex and age with the exception of an average yearly increase of 2.4% in 18–44 year-olds: IRR 1.024 (1.014–1.034). In 2014, the prevalence of AF in the adult population was 3.4%.

Conclusions We found overall stable IRs of AF for the adult Norwegian population from 2004 to 2014. The prevalence of AF was 3.4% at the end of 2014, which is higher than reported in previous studies. Signs of an increasing incidence of early-onset AF (<45 years) are worrying and need further investigation.

to 2000 and then stabilised.^{3,4} In the Framingham Heart Study, incidence rates of AF (detected by all health data sources) increased from 1958 to 2007.⁵ In Denmark, the incidence of AF causing hospitalisation increased between 1983 and 2012,⁶ while in Iceland, incidence rates of AF (inpatients and outpatients) increased among women, but not men between 1991 and 2008.⁷ A pooled estimate of age-adjusted and sex-adjusted AF prevalence reported in population studies for the period 1968–2010 suggested a prevalence of 2.8% in the adult population. However, the estimates varied from 0.6% to 6.6%, reflecting differences in the age distribution of the study populations: from 20 years and older to 65 years and older.⁸

We aimed to study trends in incidence of AF in Norway during 2004–2014, and to estimate the prevalence of AF, by age and sex.

METHODS

Data sources and data linkages

We used nationwide data on hospital visits and causes of death from the Cardiovascular Disease in Norway (CVDNOR) project.⁹ The data included time of admission and primary and secondary discharge diagnoses for all somatic inpatient hospital stays recorded in electronic patient administrative systems at all Norwegian hospitals during 1994–2009 and in the Norwegian Patient Registry from 2008 to 2014. The data also included visits to hospital outpatient clinics and to physician specialist practices with reimbursement contracts with the public health authorities registered in the Norwegian Patient Registry (2008–2014). These data were linked with data from the Norwegian Cause of Death Registry (1994–2014) and data on vital status (alive, dead or emigrated) on 1 January each year from Statistics Norway (1994–2014). The linkage was performed by Statistics Norway, using the national personal identification number.

This research was done without patient involvement.

Identification of incident and prevalent AF cases

AF was defined by the International Classification of Diseases, 9th Revision (ICD-9) diagnosis code 427.3 or ICD-10 diagnosis code I48 (including subcategories): AF and atrial flutter. AF cases thus included a minor proportion of atrial flutter

INTRODUCTION

Atrial fibrillation (AF) is an important prognostic indicator for stroke, heart failure, cardiovascular and all-cause mortality.¹ The high mortality of patients with AF highlights the importance of disease surveillance for better healthcare resource allocation and for planning of preventive public health activities.

The reported incidence and prevalence of AF have been inconsistent across studies and countries, reflecting methodological differences making comparisons difficult.² However, the majority of studies provide evidence of an increasing incidence of AF in Western societies. In Olmsted County (Minnesota, USA), incidence rates of AF (inpatients and outpatients) increased from 1980



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cases.^{10 11} The cohort of AF cases was identified from all inpatient admissions with AF as primary or secondary diagnosis during 1994–2014 and from out-of-hospital deaths with AF as underlying cause, in the adult population ≥ 18 years.

Incident AF was defined as the first inpatient admission with AF or out-of-hospital death due to AF with no previous inpatient admission with AF noted in the past 10 years. For the incidence analyses, we defined inpatient admissions with a primary or secondary diagnosis of AF as *hospitalised AF*, and AF as underlying cause of death in the Cause of Death Registry without a registered hospitalisation for AF as *out-of-hospital AF death*. A 10-year washout period was applied to minimise misclassification of recurrent cases as incident cases, while still allowing for a reasonably long study period. By this definition, less than 5% of incident cases in 2014 were recurrent cases, that is, they had received an inpatient AF diagnosis before 2014, but more than 10 years ago. Corresponding numbers using 6-year, 7-year, 8-year and 9-year washout periods were 15%, 11%, 9% and 7%, respectively.

In an analysis of the cumulative prevalence of AF from 1994 to 2014, prevalent AF cases (primary or secondary diagnoses) were identified from inpatient admissions (1994–2014) and from visits to outpatient hospital clinics and to physician specialist practices (2008–2014). Prevalent AF cases at the end of 2014 were defined as the total number of patients with at least one registered diagnosis of AF at an inpatient admission or outpatient visit during 1994–2014 and alive as of 31 December 2014.

Study design

This was a nationwide cohort study. In analyses of AF incidence, the number of yearly incident cases of inpatient admission and out-of-hospital death from AF during 2004–2014 was the numerator (online supplementary figure 1). The Norwegian population free of AF (no inpatient admission for AF in the previous 10 years) and aged ≥ 18 years on 1 January each year was the denominator. In the prevalence analysis, the number of prevalent AF cases alive as of 31 December 2014 was the numerator (online supplementary figure 2). The number of Norwegian residents aged ≥ 18 years as of 1 January 2015 was the denominator.

Comorbidity

In a supplementary analysis of inpatient admissions with incident AF, we assessed selected diagnoses co-occurring during the same admission and calculated the Charlson Comorbidity Index (CCI) using the coding algorithm and scoring suggested by Quan *et al* (online supplementary table 1).¹² We selected diagnoses based on predisposing and reported comorbid conditions for AF and comorbidity groups from the CCI.^{6 12 13} In addition, we included the diagnosis of pneumonia and markers of frailty.

Sensitivity analysis

We assessed the stability over time in incident AF cases presenting at the hospital outpatient clinic as a proportion of all incident AF cases identified, during the period with available outpatient data from 2008 to 2014. Incident hospital outpatient AF cases were identified by the first visit to a hospital outpatient clinic with a diagnosis of AF and no inpatient admissions for AF in the previous 10 years.

Statistical methods

Age-standardised incidence rates with 95% confidence intervals (CI) were calculated using direct standardisation with the age distribution of the Norwegian population free of AF as

of 1 January 2004 (ie, individuals with no hospitalisation for AF during the previous 10 years) as standard population. Age-adjusted incidence rate ratios with 95% CI were estimated, using *Poisson* regression analyses with calendar year as a continuous covariate, expressed as average yearly change in incidence rates. Average yearly change in proportion with comorbidity among incident AF cases were estimated by risk ratios with 95% CI using binomial regression analyses with calendar year as continuous covariate. An interaction term between calendar year and sex was introduced to test potential sex-differences in time trends. Risk of comorbidity in men relative to women was assessed as risk ratios with 95% CI using age-adjusted binomial regression with sex as exposure variable. In addition to overall results, we report results stratified by sex and by age groups (18–44, 45–64, 65–84 and ≥ 85 years). Stata V.15.0 (Stata, College Station, Texas, USA) software was used for all analyses.

RESULTS

Study population

We identified 175 979 cases with incident AF in Norway from 2004 to 2014 (online supplementary figure 1). Of these, 30.0% had AF as the primary diagnosis at inpatient admission, 69.4% had AF as a secondary diagnosis at inpatient admission and 0.6% were out-of-hospital AF death cases. The proportion of incident cases with AF as the primary hospital discharge diagnosis increased from 2004 (27%) to 2014 (33%) overall and across all age and sex strata (online supplementary table 2).

Women accounted for 45% of the cohort of incident AF cases and had a higher mean age \pm standard deviation at diagnosis (79.1 \pm 11.2) than men (72.1 \pm 13.0), both for primary and secondary AF diagnosis and over time (online supplementary table 3). Only 2% of the patients were younger than 45 years of age at diagnosis, 17% were 45–64, 55% were 65–84 and 26% were 85 years of age or older. The proportion of men was higher in all age groups except for the oldest group.

Comorbidity

Among hospitalised patients with AF as the primary diagnosis, 12% had CCI score of ≥ 2 in contrast to 38% among those with AF as a secondary diagnosis. The most prevalent comorbid conditions at incident inpatient admission for AF were hypertension (28%), coronary heart disease (25%), heart failure (21%) and pneumonia (13%). The majority of comorbidities were more common in men, especially co-occurring malignancy, renal disease and peripheral vascular disease. The exception was thyroid disorders, autoimmune diseases, hypertension, valvular heart disease, stroke, thromboembolism, dementia and frailty, which were more often present in women (online supplementary tables 4 and 5). The proportion with a CCI score ≥ 2 declined from 31% in 2004 to 28% in 2014 for all incident AF cases. A similar pattern was found for myocardial infarction, heart failure and diabetes, with somewhat more favourable decline over time in women than in men. For patients with AF as primary diagnosis, the proportion with co-occurring myocardial infarction increased and the proportion with heart failure was stable over time. In contrast, the co-occurrence of these conditions declined for patients with AF as secondary diagnosis (figure 1, online supplementary table 6).

Incidence

The age-standardised incidence rate of inpatient admission (primary or secondary diagnoses) or death from AF per 100 000 person years was stable, at 433 (95% CI 426 to 440) in 2004

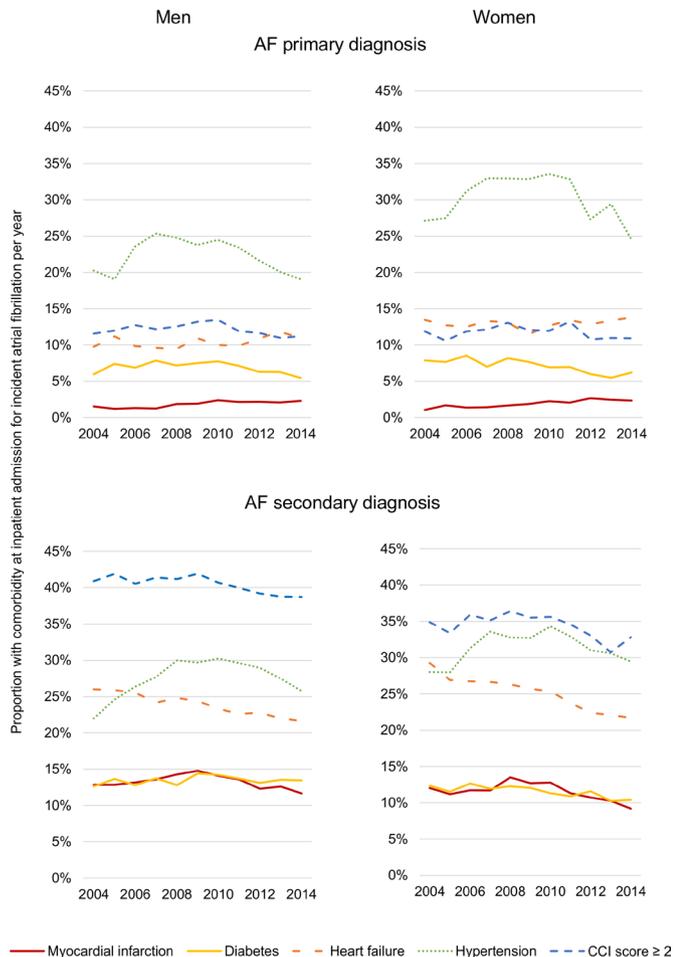


Figure 1 Proportions (%) with co-occurring myocardial infarction, diabetes, heart failure, hypertension and CCI score ≥ 2 reported at inpatient admission for incident AF from 2004 to 2014 in Norway, by year, sex and primary or secondary AF diagnosis. AF, atrial fibrillation; CCI, Charlson Comorbidity Index.

and 440 (95% CI 433 to 447) in 2014 (figure 2, table 1). On average, the total yearly incidence rate did not change markedly (-0.2% , 95% CI -0.3% to 0.0% , $p=0.02$). Consistent rates across all strata of age group and sex were observed with the

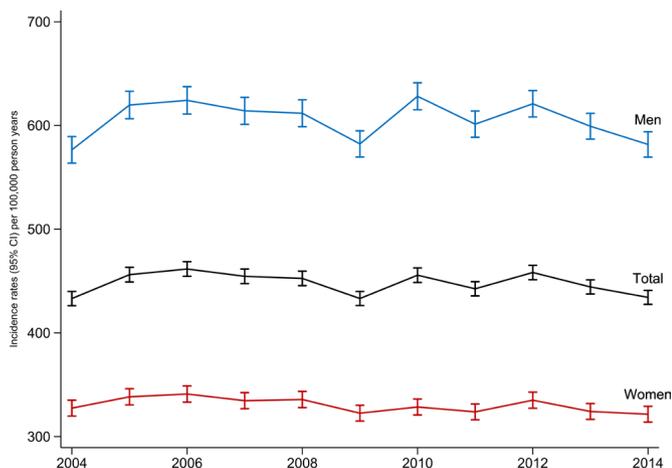


Figure 2 Age-standardised incidence rates with 95% CI of atrial fibrillation per 100 000 person years from 2004 to 2014 in Norway by year and sex.

exception of those aged 18–44 years where we observed a 2.4% (95% CI 1.4% to 3.4%, $p<0.001$) increase in the overall yearly incidence rate (women 2.1% (95% CI -0.1% to 4.3%, $p=0.06$), men 2.4% (95% CI 1.1% to 3.5%, $p<0.001$)) (figure 3, table 2).

The age-standardised incidence rate of inpatient admission with a primary diagnosis of AF per 100 000 person years increased from 117 (95% CI 113 to 120) in 2004 to 142 (95% CI 139 to 146) in 2014 (online supplementary figure 3 and table 3). The overall trend was a mean yearly increase of 1.9% (95% CI 1.7% to 2.2%, $p<0.001$). The increase was consistent in women (2.1% per year, 95% CI 1.7% to 2.6%, $p<0.001$) and men (1.6% per year, 95% CI 1.3% to 2.0%, $p<0.001$) and across age groups in both sexes, although the increase was small among those aged 45–64 years (online supplementary figure 4 and table 7).

In analysis of incident cases with AF as a secondary diagnosis at inpatient admission, we observed a stable to slightly decreasing trend in the incidence rate per 100 000 person years, from 313 (95% CI 308 to 319) in 2004 to 295 (95% CI 290 to 301) in 2014 (online supplementary figure 5 and table 3). On average, the yearly decrease was 1.1% (95% CI 0.9% to 1.2%, $p<0.001$). This finding was consistent across sex and age groups, even though the estimates for those aged 18–45 years were imprecise (online supplementary figure 6 and table 7).

Cumulative prevalence

Since 1994, AF had been diagnosed during an inpatient admission or outpatient visit in 136 828 individuals (women: 55 440 (41%); men: 81 388 (59%)) in Norway as of 31 December 2014. This corresponds to a cumulative prevalence of 3.4% of the total Norwegian population ≥ 18 years (2.8% in women and 4.0% in men) at the end of 2014. Figure 4 and table 3 show the prevalence by age for men and women.

Sensitivity analysis of outpatient visits

In total, 77 494 patients were registered with an AF diagnosis at one or more visits to a hospital outpatient clinic during 2008–2014. The number of outpatients with no inpatient admission for AF in the preceding 10 years totalled 30 185, increasing from 4041 in 2008 to 4895 in 2014, corresponding to 20% and 23% of all incident AF cases.

DISCUSSION

We observed stable age-standardised incidence rates of AF from 2004 to 2014. The findings were generally consistent across sex and age groups except for a mean yearly increase of 2.4% in the youngest age group (18–44 years), which was most prominent among men. The cumulative prevalence of AF was 3.4% in the adult Norwegian population.

Temporal changes in coding practice, reimbursement rates, clinical guidelines, treatment strategies and handling of patients have to be taken into account when interpreting registry data from patient administrative systems. These changes may influence the preference of AF as primary or secondary diagnosis and could partly explain the decreased incidence of hospitalisations with AF as a secondary diagnosis. Alternatively, the apparent decrease reflects the declining incidence rates of associated conditions such as acute myocardial infarction and heart failure in Norway.^{14 15} Comorbidity burden and proportions with co-occurring myocardial infarction and heart failure for all incident AF cases in our study decreased over time, as previously shown.^{4–6 16–18} However, two population-based health examination surveys found stable rates of comorbid myocardial infarction,

Arrhythmias and sudden death

Table 1 Age-standardised incidence rates of inpatient admission or death from atrial fibrillation per 100 000 person years in Norway 2004–2014 by year, sex and age group

	2004	2005	2006	2007	2008	2009	2010	2011	2012	2013	2014
<i>Men</i>											
18–44 years											
Incidence rate	26	31	31	33	36	30	29	36	36	37	35
Cases	225	267	268	287	322	275	264	330	339	346	335
45–64 years											
Incidence rate	306	314	336	323	316	302	320	313	316	328	291
Cases	1696	1797	1983	1987	2000	1958	2069	2044	2086	2170	1939
65–84 years											
Incidence rate	2147	2238	2249	2261	2187	2073	2231	2158	2286	2158	2125
Cases	4774	4986	4963	4986	4851	4657	5079	4993	5455	5273	5335
≥85 years											
Incidence rate	5484	6531	6353	5956	6413	6207	6436	6232	6068	6006	6195
Cases	1192	1458	1468	1421	1548	1536	1637	1600	1603	1596	1691
All ages											
Incidence rate	576	620	624	614	612	582	617	602	620	602	589
Cases	7887	8508	8682	8681	8721	8426	9049	8967	9483	9385	9300
<i>Women</i>											
18–44 years											
Incidence rate	7	8	8	8	9	10	9	9	11	8	9
Cases	61	71	69	68	77	84	81	77	100	74	84
45–64 years											
Incidence rate	108	118	108	115	120	105	106	104	109	109	105
Cases	601	675	648	710	766	685	701	694	717	717	695
65–84 years											
Incidence rate	1241	1244	1248	1226	1235	1205	1204	1220	1280	1212	1231
Cases	3996	3975	3908	3772	3750	3633	3613	3681	3915	3748	3851
≥85 years											
Incidence rate	4063	4357	4592	4369	4263	4088	4074	4118	4042	4141	4066
Cases	2299	2487	2719	2650	2642	2575	2619	2656	2633	2673	2640
All ages											
Incidence rate	327	338	341	335	336	323	322	325	335	325	326
Cases	6957	7208	7344	7200	7235	6977	7014	7108	7365	7212	7270
<i>Total</i>											
18–44 years											
Incidence rate	17	20	20	21	23	20	19	22	24	23	23
Cases	286	338	337	355	399	359	345	407	439	420	419
45–64 years											
Incidence rate	207	216	222	219	218	204	213	209	213	219	199
Cases	2297	2472	2631	2697	2766	2643	2770	2738	2803	2887	2634
65–84 years											
Incidence rate	1620	1663	1670	1666	1643	1582	1648	1625	1717	1624	1622
Cases	8770	8961	8871	8758	8601	8290	8692	8674	9370	9021	9186
≥85 years											
Incidence rate	4458	4980	5088	4824	4866	4685	4750	4718	4636	4693	4729
Cases	3491	3945	4187	4071	4190	4111	4256	4256	4236	4269	4331
All ages											
Incidence rate	433	456	462	455	453	433	447	443	458	446	440
Cases	14844	15716	16026	15881	15956	15403	16063	16075	16848	16597	16570

Incidence rates including 95% CIs are illustrated in figures 2 and 3.

more in agreement with the novel finding from our study of increasing proportions of co-occurring myocardial infarction for admissions with AF as primary diagnosis.^{4 5} In contrast to our study, a Danish registry study reported lower, but increasing levels of comorbidity burden over time.⁶ Of interest, we found higher comorbidity in men with more co-occurring malignancy, renal disease and peripheral vascular disease, whereas women

who were older had more often dementia, frailty and autoimmune disease. We also found that men had more co-occurring coronary heart disease and women more hypertension, thyroid disorders and stroke, similar to previous reports.^{5 16}

Detection of AF has possibly increased during the study period due to increased awareness of AF and its complications among clinicians and in the population.^{1 5} Furthermore, it has been

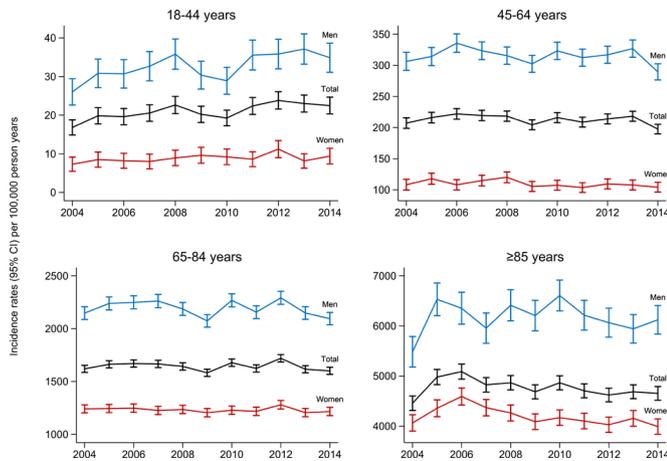


Figure 3 Age-standardised incidence rates with 95% CI of atrial fibrillation per 100 000 person years from 2004 to 2014 in Norway by year, sex and age group.

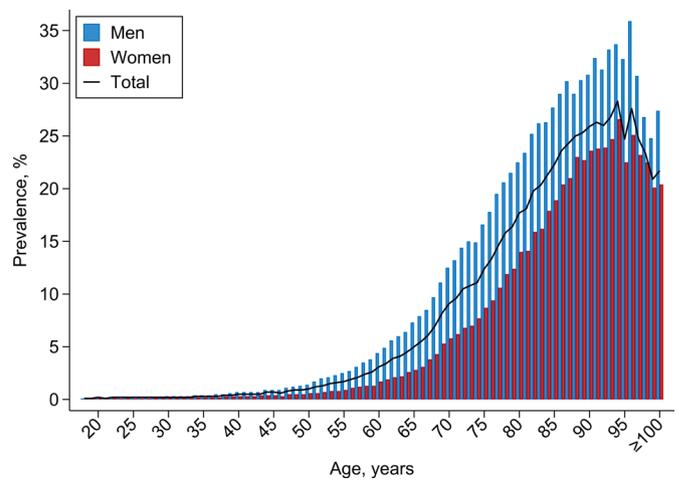


Figure 4 Cumulative prevalence of atrial fibrillation from 1994 to 2014 in Norway by age.

suggested that increased incidence rates may partly arise due to improvements in surveillance methods (new modalities for short-term and long-term ECG monitoring) and their increased use in different patient groups (eg, poststroke ECG screening). In the population-based Framingham Heart Study, incidence rates of AF detected by all health data sources increased from 1958 to 2007, yet the rate of AF detected by ECG at repeated study visits did not increase during this period.⁵

Our results suggested a shift over time in referral practice to more outpatient treatment for new patients with AF, similar to reports from a Danish registry study.⁶ The described stable incidence rates of AF may therefore have been increasingly underestimated and likely reflect an unbiased upward trend which supports the observation that incidence rates have not decreased.

Changes in AF incidence over time are of clinical interest and need surveillance. Our observed increased incidence rate of AF in young adults is worrisome and requires further study of changes

in underlying risk factors, primarily factors that impact younger birth cohorts born after the 1950s–1960s differently than older cohorts. In Norway, prevalence of obesity has increased over time, especially in the young adult population.¹⁹ In addition, the non-declining incidence rates of acute myocardial infarction in those below 45 years during 2001–2014 in Norway, could add to AF risk.¹⁴ Nevertheless, declining blood pressure has been observed in younger cohorts, which may lower the AF risk.²⁰ Increased awareness of obesity as an important modifiable risk factor in younger cohorts, as well as preventive measures that promote a stable or reduced body mass index over time, may reduce AF incidence.²¹

The reported incidence of AF has been inconsistent across published studies, depending on approaches used. In different US health registry databases, AF incidence rates increased from 2001 to 2008²² and from 2004 to 2016.²³ In line with our finding of a stable incidence from 2004 to 2014, AF incidence rates increased from 1998 up to 2007 and then plateaued from 2007 to 2010 in a UK primary care database linked with hospitalisation data.¹⁷ Incidence rates of AF in the Korean National Health Insurance database were stable from 2006 to 2015.²⁴ In a Western Australian population, incidence rates of hospitalisations with AF as the primary diagnosis increased, whereas incidence rates of hospitalisations with AF as any diagnosis decreased from 1995 to 2010, using a hospital admission database,¹⁸ in line with our findings.

In a systematic review, Ball *et al*⁸ estimated a pooled prevalence of 2.8% (95% CI 2.3% to 3.4%) in adults, 2.4% (95% CI 1.9% to 2.9%) in females and 3.3% (95% CI 2.7% to 4.0%) in males. Interestingly, we found a prevalence in Norway at the upper limit of these CIs, namely 3.4% in adults, 2.8% in women and 4.0% in men. To our knowledge, this is the highest nationwide prevalence reported to date. Based on inpatient and outpatient hospital records, the prevalence was 2.9% in the Swedish adult population in 2010.²⁵ A similar prevalence, 3.0%, was found in a smaller cohort in Northern Sweden in 2010, using hospital records and a regional ECG database.²⁶

A major strength of our study is the use of nationwide data, and a long observation period, which in turn allow for an adequate washout period and enabled calculations of temporal trends of incidence. Our analysis of AF incidence is limited to inpatient admissions only as data from outpatient visits did not allow for an adequate washout period to define incident cases and were further limited by poor reporting: the proportion of

Table 2 Average yearly change in incidence rates of inpatient admission or death from atrial fibrillation in Norway 2004–2014 by sex and age group

	Cases	Incidence rate ratio (95% CI)	P value
Men			
18–44 years	3258	1.024 (1.013 to 1.035)	<0.001
45–64 years	21 729	0.996 (0.992 to 1.000)	0.062
65–84 years	55 352	0.998 (0.995 to 1.001)	0.117
≥85 years	16 750	1.000 (0.995 to 1.005)	0.987
All ages	97 089	0.999 (0.997 to 1.001)	0.486
Women			
18–44 years	846	1.021 (0.999 to 1.043)	0.060
45–64 years	7609	0.991 (0.984 to 0.998)	0.016
65–84 years	41 842	0.999 (0.996 to 1.002)	0.439
≥85 years	28 593	0.995 (0.991 to 0.998)	0.004
All ages	78 890	0.994 (0.992 to 0.997)	<0.001
Total			
18–44 years	4104	1.024 (1.014 to 1.034)	<0.001
45–64 years	29 338	0.995 (0.991 to 0.999)	0.009
65–84 years	97 194	0.999 (0.998 to 1.001)	0.611
≥85 years	45 343	0.998 (0.995 to 1.001)	0.132
All ages	175 979	0.998 (0.997 to 1.000)	0.020

Table 3 Cumulative prevalence of atrial fibrillation from 1994 to 2014 in Norway by age

Age group, years	Men			Women			Total		
	Cases	Population	Prevalence, %	Cases	Population	Prevalence, %	Cases	Population	Prevalence, %
18–24	356	245 638	0.1	311	231 734	0.1	667	477 372	0.1
25–29	413	179 774	0.2	253	173 546	0.1	666	353 320	0.2
30–34	552	178 443	0.3	281	167 907	0.2	833	346 350	0.2
35–39	841	176 828	0.5	420	165 215	0.3	1261	342 043	0.4
40–44	1422	192 531	0.7	618	181 192	0.3	2040	373 723	0.5
45–49	2112	192 502	1.1	799	180 774	0.4	2911	373 276	0.8
50–54	3257	172 907	1.9	1154	163 707	0.7	4411	336 614	1.3
55–59	4996	160 450	3.1	1782	155 254	1.1	6778	315 704	2.1
60–64	7889	144 730	5.5	2944	142 764	2.1	10833	287 494	3.8
65–69	12 292	138 419	8.9	5359	139 283	3.8	17 651	277 702	6.4
70–74	13 169	95 105	13.8	6734	101 758	6.6	19 903	196 863	10.1
75–79	12 000	63 154	19.0	7987	76 146	10.5	19 987	139 300	14.3
80–84	10 830	43 980	24.6	9588	61 543	15.6	20 418	105 523	19.3
85–89	7534	25 875	29.1	9577	45 535	21.0	17 111	71 410	24.0
90–94	3158	9881	32.0	6082	25 043	24.3	9240	34 924	26.5
95–99	524	1631	32.1	1402	6063	23.1	1926	7694	25.0
≥100	43	157	27.4	149	729	20.4	192	886	21.7
All ages	81 388	2 022 005	4.0	55 440	2 018 193	2.7	136 828	4 040 198	3.4

specialists that reported their activity to the Norwegian Patient Registry was 56% in 2009.²⁷ In addition, our definition of AF cases was based on inpatient hospital registry data and not verified by ECG or additional medical records. However, the validity of AF reported in Scandinavian studies has been high. Among patients with a hospital register-based AF diagnosis code, AF was confirmed by medical records or ECG from hospitals or primary care in 89%–97%.^{11 28 29} Another limitation is that we lacked data from primary care physicians and nursing homes to better estimate the true prevalence of AF. While we do not know the proportion of patients with AF in Norway that are treated only out-of-hospital, a study by Friberg *et al*²⁵ suggested that 22% of Swedish patients with AF were seen only in primary care practices. If we extrapolate this to our results, the prevalence of AF in the adult Norwegian population would be as high as 4.1%.

CONCLUSION

For the adult population in Norway, we found overall stable incidence rates of AF from 2004 to 2014 and an AF prevalence of at

least 3.4% at the end of 2014. Signs of increased incidence rates of early-onset AF (<45 years) are of concern and need further investigation.

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Contributors GST, IA, JI and RS planned the work. LJK, JI and IA had access to the data and contributed to the statistical analyses. LJK drafted the manuscript. All authors contributed to the design of the study, have critically revised the draft and have approved the final version of the submitted manuscript. LJK and IA are responsible for the overall content as guarantors.

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Competing interests MM reports personal fees from Boehringer-Ingelheim, personal fees from Bayer, personal fees from Bristol-Myers Squibb, personal fees from MSD, outside the submitted work. TB reports personal fees from Boehringer-Ingelheim, Bayer and Pfizer/Bristol-Myers Squibb, outside the submitted work. IEC reports personal fees from Merck, outside the submitted work. The other authors have nothing to disclose.

Patient and public involvement Patients and/or the public were not involved in the design, or conduct, or reporting, or dissemination plans of this research.

Patient consent for publication Not required.

Key messages

What is already known on this subject?

- ▶ The reported incidence and prevalence of atrial fibrillation have been inconsistent across studies and countries.

What might this study add?

- ▶ The incidence rates of atrial fibrillation among adults in Norway were stable from 2004 to 2014, except for a possible increase in those less than 45 years. The prevalence of atrial fibrillation was 3.4% at the end of 2014.

How might this impact on clinical practice?

- ▶ A possible increase in early-onset atrial fibrillation incidence warrants increased awareness towards modifiable risk factors.

Ethics approval The study was approved by the Regional Committee for Medical and Health Research Ethics, Health Region West (on 6 October 2009; reference number 2009/861) and complies with the Declaration of Helsinki.

Provenance and peer review Not commissioned; externally peer reviewed.

Data availability statement No data are available. The approval from the Ethics Committee does not include permission to make data materials available. LJK, JI and IA had full access to all the data in the study and takes responsibility for its integrity and the data analysis.

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