Original Article

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Association Between Atrial Fibrillation and the Risk of Dementia in the Korean Elderly: A 10-Year Nationwide Cohort Study

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Objectives: The objective of this study was to determine the effect of atrial fibrillation (AF) on the risk of dementia in the Korean elderly.

Methods: A 10-year retrospective cohort study was conducted using the National Health Insurance Service-Senior Cohort database. We excluded those who were under 65 years of age as of January 2006 (n=46113), those who were diagnosed with dementia between 2002 and 2005 (n=9086), and those with a history of stroke prior to AF diagnosis (n=8392). We used a Cox proportional hazards model with a time-varying covariate to determine whether AF is associated with the risk of dementia after adjusting for potential confounders.

Results: In univariable Cox regression, the hazard ratio (HR) of dementia according to AF status was 1.28 (95% confidence interval [CI], 1.23 to 1.33). After adjusting for potential confounders, AF was found to increase the risk of dementia (HR, 1.12; 95% CI, 1.07 to 1.17), Alzheimer dementia (HR, 1.12; 95% CI, 1.07 to 1.17), and vascular dementia (HR, 1.10; 95% CI, 1.03 to 1.18). In patients diagnosed with AF, the incidence of dementia was lower (HR, 0.50; 95% CI, 0.47 to 0.52) in patients who were treated with oral anticoagulants.

Conclusions: Investigating the potential risk factors of dementia in an aged society is important. We found a slightly higher risk of de-

mentia in those with AF than in those without AF, and we therefore concluded that AF is a potential risk factor for dementia.

Key words: Dementia, Atrial fibrillation, Risk factors, Primary prevention

INTRODUCTION

Korea became an aged society in 2017, with elderly citizens (aged 65 and up) accounting for over 14% of the population.

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The prevalence of dementia, a disorder predominantly affecting elderly individuals, is also on the rise as the population ages. More than 10% of those aged 65 and over had dementia in 2017, and the number of patients with dementia is expected to surpass 1 million nationwide by 2024 [1]. The level of socioeconomic burden posed by the disease also continues to increase, with the cost of managing dementia projected to reach 18.8 trillion Korean won by 2020 [1]. Against this backdrop, it is important to prioritize the development of primary preventive policies in addition to ways to manage existing patients; in the service of this goal, further studies are necessary to clarify the potential risk factors for dementia.

Risk factors for dementia include socio-demographic fac-

tors, such as age and sex; genetic factors; physical and mental health factors, such as hypertension, diabetes, and depression; and lifestyle habits, such as smoking and drinking [2-4]. Many studies of community-based cohorts in Europe and the United States have found that atrial fibrillation (AF) is an independent risk factor for dementia [5-11], while some studies have reported that AF increases not only the risk of vascular dementia, but also that of Alzheimer dementia.

AF is the most common type of chronic arrhythmia and shares risk factors with dementia, including age, hypertension, diabetes, and smoking. Additionally, AF is a risk factor for ischemic stroke, which is known to cause vascular dementia [12]. Studies are being conducted to understand the mechanisms by which AF independently causes dementia [13], with hypotheses including asymptomatic cerebral infarction [14], chronic cerebral hypoperfusion [15], inflammation [16,17], decreased brain volume [18], and microbleeds due to anticoagulant use. Meanwhile, other studies have claimed that treatment with catheter ablation [19] as well as the use of oral coagulants in AF patients can actually reduce the risk of dementia [20]. Further studies are needed to assess the association between the use of coagulants and the occurrence of dementia.

Outside of Korea, many studies have already probed the association between AF and dementia. However, this has not

been the case in Asian countries, including Korea, despite the fact that the epidemiological characteristics of AF and dementia varies by country and ethnic group [21-25]. AF and dementia are chronic diseases that are affected by lifestyle habits [26-28] and genetic factors [29-31], so they are affected by racial and cultural factors. Therefore, it is crucial to study this topic in the Korean population, rather than simply applying the results of studies conducted abroad.

For this study, we used the National Health Insurance Service-Senior Cohort (NHIS-SC) database, which accurately represents the nation's population and has a large sample size, to assess whether AF acts as an independent risk factor for dementia, Alzheimer dementia, and vascular dementia. And we also examined the impact of anticoagulant use on the dementia risk of AF patients.

METHODS

Data Source and Study Population

This study analyzed data from the NHIS-SC. The NHIS-SC database consists of 14 years (2002-2015) of cumulative records of 550 000 randomly-selected members of the senior population aged 60 years and older more regarding their use of medical services. This sample data accounts for approximately 10%

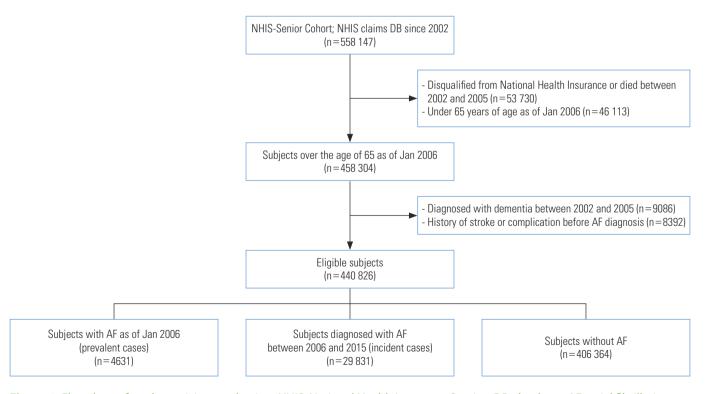


Figure 1. Flowchart of study participant selection. NHIS, National Health Insurance Service; DB, database; AF, atrial fibrillation.

of the 5 581 470-member senior population of Korea. The personal information of the individuals was not identifiable.

The study determined the washout period of dementia to be 4 years; therefore, of the 14 years of follow-up data, we analyzed dementia risk over the 10-year period from 2006 to 2015. From the total cohort of 558 147 seniors, those who were not eligible for National Health Insurance or Medical Aid as of January 2006 ($n=53\,730$) for reasons such as death and those under 65 years of age ($n=46\,113$) were excluded. A final sample of 440 826 subjects were included in the analysis after excluding subjects who had been diagnosed with dementia between 2002 and 2005 (n=9086) and subjects with a history of stroke prior to diagnosis of AF (n=8392) (Figure 1).

Operational Definitions of Diseases

Cases of dementia, AF, and other comorbidities were defined as cases with the disease in question listed as the main diagnosis or sub-diagnosis based on the Korean Standard Classification of Diseases. The code for AF (the main independent variable) was I48, the codes for dementia were F00-F03, and the codes for Alzheimer and vascular dementia were F00 and F01, respectively. The codes for hypertension were I10-I15, the codes for diabetes were E10-E14, those for stroke were I60-I63, those for post-stroke symptoms were I690-I694, the code for heart failure was I50, the code for ischemic heart disease was I24, and the codes for valvular heart diseases were I05-I08 and I34-I37. Stroke was limited to cases for which the insurance claim was classified as inpatient and magnetic resonance imaging or computed tomography examination was performed.

Anticoagulant use was defined as the prescription of drugs that contained the main ingredient codes for wafarin and direct oral anticoagulants. The main ingredient codes were 249101ATB, 249102ATB, 249103ATB, 249104ATB, 249105ATB, 249106ATB, 249107ATB, 249108ATB, 249109ATB, 511401ATB, 511402ATB, 511403ATB, 511404ATB, 613701ACH, 613702ACH, 613703ACH, 617001ATB, 617002ATB, 643601ATB, 643602ATB, and 643603ATB.

Statistical Analysis

The general characteristics of the subjects were identified using the *t*-test and the chi-square test. A Cox proportional hazards model was used with the independent variable of AF status as a time-varying covariate over the 10-year follow-up study period to explore the association between AF and dementia. The adjusted confounders were age, sex, hyperten-

sion, diabetes, heart failure, ischemic heart disease, and valvular heart disease. The same method was used to assess whether AF acted as an independent risk factor for Alzheimer and vascular dementia, as well as whether anticoagulant use affected the development of dementia among AF patients.

All statistical analyses were performed with SAS version 9.4 (SAS Institute Inc., Cary, NC, USA).

Ethics Statement

This study was approved by the Institutional Review Board of Yeungnam University Hospital (YUMC-2018-08-009).

RESULTS

Of the total of 440 826 subjects, 4631 (1.1%) patients had already been diagnosed with AF as of January 2006, a total of 29 831 (6.8%) patients were newly diagnosed with AF between 2006 and 2015, and 406 364 (92.2%) subjects were not diagnosed with AF (Table 1).

Table 1. Baseline demographic and clinical characteristics of subjects

| | Total | Atrial fibrillation | | | | |
|------------------------|--------------|---------------------|--------------------------|-----------------|--|--|
| Variables | Total (n) | Prevalent cases | Incident cases (10 y) | No diagnosis | | |
| Age (mean \pm SD, y) | 72.8±6.5 | 74.1 ± 6.7 | 73.4 ± 6.3 | 72.7 ± 6.5 | | |
| Sex | | | | | | |
| Male | 176 330 | 2089 (1.2) | 13 187 (7.5) | 161 055 (91.3) | | |
| Female | 264 496 | 2542 (1.0) | 16 644 (6.3) | 245 309 (92.7) | | |
| Hypertension | | | | | | |
| Yes | 184 706 | 3621 (2.0) | 14 677 (7.9) | 166 408 (90.1) | | |
| No | 256 120 | 1010 (0.4) | 15 154 (5.9) | 239 956 (93.7) | | |
| Diabetes mellitus | | | | | | |
| Yes | 92 133 | 1903 (2.1) | 6835 (7.4) | 83 395 (90.5) | | |
| No | 348 693 | 2728 (0.8) | 22 996 (6.6) | 322 970 (92.6) | | |
| Heart failure | | | | | | |
| Yes | 13 666 | 1385 (10.1) | 1726 (12.6) | 10 555 (77.2) | | |
| No | 427 160 | 3246 (0.8) | 28 105 (6.6) | 395 809 (92.6) | | |
| Ischemic heart disea | ase | | | | | |
| Yes | 1763 | 111 (6.3) | 170 (9.6) | 1482 (84.1) | | |
| No | 439 063 | 4520 (1.0) | 29 661 (6.8) | 404 882 (92.2) | | |
| Valvular heart disea | se | | | | | |
| Yes | 3086 | 468 (15.2) | 460 (14.9) | 2158 (69.9) | | |
| No | 437 740 | 4163 (1.0) | 29 371 (6.7) | 404 206 (92.3) | | |
| Total | 440 826 | 4631 (1.1) | 29 831 (6.8) | 406 364 (92.2) | | |

Values are presented as number (%).

SD, standard deviation.

Table 2. Univariable and multivariable Cox regression for the incidence of dementia, Alzheimer dementia, and vascular dementia

| | | Dementia | | | Alzheimer dementia | | Vascular dementia | |
|------------------------|---------------------------------|-----------------|--|-----------------|----------------------------------|-----------------|--|-----------------|
| Variables | Univariable Cox (unadjusted) | <i>p</i> -value | Multivariable Cox (adjusted) ¹ | <i>p</i> -value | Multivariable Cox (adjusted)¹ | <i>p</i> -value | Multivariable Cox (adjusted) ¹ | <i>p</i> -value |
| Age | 1.08 (1.08, 1.09) | < 0.001 | 1.08 (1.08, 1.08) | < 0.001 | 1.09 (1.09, 1.09) | < 0.001 | 1.08 (1.08, 1.08) | < 0.001 |
| Sex | | | | | | | | |
| Male | 1.00 (reference) | | 1.00 (reference) | | 1.00 (reference) | | 1.00 (reference) | |
| Female | 1.47 (1.45, 1.49) | < 0.001 | 1.26 (1.25, 1.28) | < 0.001 | 1.31 (1.29, 1.32) | < 0.001 | 1.08 (1.06, 1.11) | < 0.001 |
| Hypertension | | | | | | | | |
| No | 1.00 (reference) | | 1.00 (reference) | | 1.00 (reference) | | 1.00 (reference) | |
| Yes | 1.33 (1.31, 1.34) | < 0.001 | 1.14 (1.13, 1.16) | < 0.001 | 1.10 (1.08, 1.12) | < 0.001 | 1.36 (1.33, 1.40) | < 0.001 |
| Diabetes mellitus | | | | | | | | |
| No | 1.00 (reference) | | 1.00 (reference) | | 1.00 (reference) | | 1.00 (reference) | |
| Yes | 1.23 (1.21, 1.24) | < 0.001 | 1.23 (1.21, 1.25) | < 0.001 | 1.23 (1.21, 1.25) | < 0.001 | 1.28 (1.25, 1.31) | < 0.001 |
| Heart failure | | | | | | | | |
| No | 1.00 (reference) | | 1.00 (reference) | | 1.00 (reference) | | 1.00 (reference) | |
| Yes | 1.39 (1.36, 1.41) | < 0.001 | 1.08 (1.06, 1.10) | < 0.001 | 1.09 (1.07, 1.11) | < 0.001 | 1.09 (1.05, 1.12) | < 0.001 |
| Ischemic heart dis | ease | | | | | | | |
| No | 1.00 (reference) | | 1.00 (reference) | | 1.00 (reference) | | 1.00 (reference) | |
| Yes | 1.20 (1.18, 1.22) | < 0.001 | 1.05 (1.04, 1.07) | < 0.001 | 1.05 (1.03, 1.07) | < 0.001 | 1.07 (1.04, 1.10) | < 0.001 |
| Valvular heart dise | ease | | | | | | | |
| No | 1.00 (reference) | | 1.00 (reference) | | 1.00 (reference) | | 1.00 (reference) | |
| Yes | 1.31 (1.20, 1.43) | < 0.001 | 1.02 (0.93, 1.12) | 0.697 | 1.02 (0.92, 1.13) | 0.697 | 1.10 (0.94, 1.28) | 0.215 |
| Atrial fibrillation (t | ime-varying covariate) | | | | | | | |
| No | 1.00 (reference) | | 1.00 (reference) | | 1.00 (reference) | | 1.00 (reference) | |
| Yes | 1.28 (1.23, 1.33) | < 0.001 | 1.12 (1.07, 1.17) | < 0.001 | 1.12 (1.07, 1.17) | < 0.001 | 1.10 (1.03, 1.18) | 0.004 |

Values are presented as hazard ratio (95% confidence interval).

Adjusted for age, sex, hypertension, diabetes mellitus, heart failure, ischemic heart disease, and valvular heart disease.

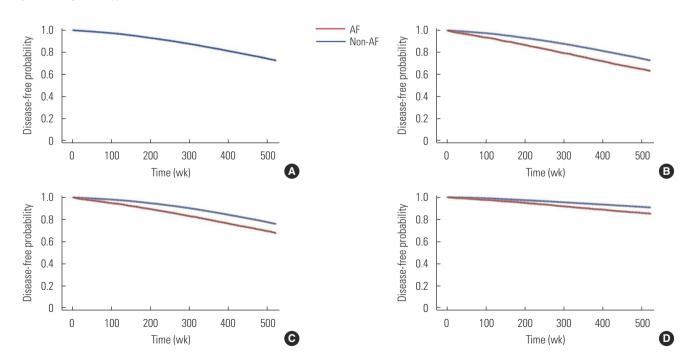


Figure 2. Kaplan-Meier curves for the incidence of dementia. Survival curve for overall incidence of (A) dementia, (B) dementia according to atrial fibrillation (AF), (C) Alzheimer's dementia according to AF, and (D) vascular dementia according to AF.

Table 3. Risk of dementia, Alzheimer dementia, and vascular dementia according to oral anticoagulant use in atrial fibrillation patients (multivariable Cox regression)

| Diseases | HR (95% CI) ¹ | <i>p</i> -value |
|--------------------|--------------------------|-----------------|
| Dementia | 0.50 (0.47, 0.52) | < 0.001 |
| Alzheimer dementia | 0.48 (0.45, 0.51) | < 0.001 |
| Vascular dementia | 0.55 (0.50, 0.60) | < 0.001 |

HR, hazard ratio; CI, confidence interval.

¹Adjusted for age, sex, hypertension, diabetes mellitus, heart failure, ischemic heart disease, and valyular heart disease.

The Cox regression hazard model with AF status as a timevarying covariate showed that age, sex, hypertension, diabetes, heart failure, ischemic heart disease, and valvular heart disease were each meaningfully associated with the risk of dementia when not adjusted for the effects across the variables. The hazard ratio (HR) of dementia according to AF status was 1.28 (95% confidence interval [CI], 1.23 to 1.33). The multivariable analysis, which adjusted for age, sex, hypertension, diabetes, heart failure, ischemic heart disease, and valvular heart disease, showed that having AF slightly increased the risk of dementia (HR, 1.12; 95% CI, 1.07 to 1.17) (Table 2, Figure 2A and 2B). When we adiusted for the same variables to assess the associations between AF and Alzheimer dementia and between AF and vascular dementia, AF was found to be significantly associated with both Alzheimer dementia (HR, 1.12; 95% CI, 1.07 to 1.17) and vascular dementia (HR, 1.10; 95% CI, 1.03 to 1.18) (Table 2, Figure 2C and 2D).

Among AF patients, those who used oral anticoagulants had a statistically significantly lower risk of developing dementia (HR, 0.50; 95% CI, 0.47 to 0.52). Anticoagulant use was also confirmed to reduce the risk of Alzheimer dementia (HR, 0.48; 95% CI, 0.45 to 0.51) and vascular dementia (HR, 0.55; 95% CI, 0.50 to 0.60) (Table 3).

DISCUSSION

This study is novel for its attempt to epidemiologically understand the potential risk factors for dementia by analyzing the association between AF and dementia. The use of National Health Insurance Service (NHIS) sample cohort data improved the sample size and representation of the subjects, while also ensuring a sufficient follow-up period. Since the NHIS sample cohort data are secondary data collected for the purpose of administrative management, it was important to clarify the standards used to define diseases and morbidity, as well as

the selection method of subjects. In this study, the washout period was set at 4 years, given that the 99th percentile of the average time to diagnosis in patients diagnosed with dementia was 38 months, to rule out subjects who had already been diagnosed with dementia. Subjects who had already been diagnosed with stroke before the diagnosis of AF were also excluded to eliminate cases of dementia caused by stroke preceding AF.

This study verified that the association between AF and dementia was statistically significant when controlling for sex, age, hypertension, diabetes, heart failure, and valvular heart disease. The association between AF and dementia has 2 explanations. First, AF and dementia share common risk factors. such as sex, age, lifestyle habits, hypertension, diabetes, and heart disease [3,4,32-34], and second, AF increases the risk of dementia through a number of potential mechanisms. The results of this study serve as the foundation for the second explanation, that AF itself may contribute to an increased risk of dementia. The results also showed that the development of dementia, Alzheimer dementia, and vascular dementia was meaningfully associated with AF status even when controlling for sex, age, and comorbidities. This aligns with preceding meta-analyses [35-37] as well as registry-based studies [6,8,37]. These results can be complemented by research on potential mechanisms by which AF may increase the risk of dementia. Suggested mechanisms include cerebral hypoperfusion due to low cardiac output [15], asymptomatic stroke due to a small thrombus [14], and activation of inflammation [17].

Many preceding studies [38,39] have found that the use of anticoagulants lowers the risk of deterioration of cognitive functions. The suspected mechanism behind this is that the use of anticoagulants can prevent blood clots that can lead to symptomatic and asymptomatic stroke [39]. This study also confirmed that the use of anticoagulants lowered the risk of developing dementia, Alzheimer's dementia, and vascular dementia. From this, it is safe to assume that the consistent use of anticoagulants in AF patients may help lower the risk of developing dementia and the associated deterioration of cognitive functions.

The subjects of this study were senior Korean citizens aged 65 and older. Given that AF and dementia are becoming increasingly prevalent among the middle-aged [21,40] and that a previous study has shown a higher risk of developing dementia among younger AF patients [5], future studies should be carried out on a broader population base that includes younger



population to verify whether AF raises the risk of dementia among these subjects as well.

Meanwhile, unlike previous studies [5,8,12,37] such as the Framingham Heart Study, which used health examination data collected for the purpose of the study, the NHIS elderly sample cohort database was originally collected for the purposed of reviewing medical claims under disease diagnosis codes. This implies limitations in accurately assessing morbidity and prevalence based on symptoms, signs, and diagnostic exam results. Surveys and diagnosis-based research should be carried out in the future to ensure a more detailed define for target diseases.

Despite the above limitations, this study attempted to move towards obtaining an epidemiological explanation of dementia among AF patients by utilizing access to the big sample data regarding the use of medical services available from the NHIS senior citizen cohort database. The results of this study confirmed that AF can potentially increase the risk of dementia, Alzheimer dementia, and vascular dementia, and that the intake of oral anticoagulants lowers the risk of developing dementia among AF patients in Korea.

CONFLICT OF INTEREST

The authors have no conflicts of interest associated with the material presented in the paper.

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AUTHOR CONTRIBUTIONS

Conceptualization: MAN, KSL, TYH. Data curation: MAN. Formal analysis: MAN. Funding acquisition: None. Methodology:

MAN, KSL. Project administration: KSL. Visualization: MAN. Writing-original draft: MAN, KSL, TYH. Writing - review & editing: MAN, KSL, TYH.

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