SCIENTIFIC LETTER

Arterial oxygen desaturation during sleep and atrial fibrillation

T Tanigawa, K Yamagishi, S Sakurai, I Muraki, H Noda, T Shimamoto, H Iso

Heart 2006;92:1854-1855. doi: 10.1136/hrt.2005.081257

trial fibrillation is a common arrhythmia with associated complications of stroke and other adverse outcomes. We conducted a population-based study of 1763 Japanese men aged 40–74 years to examine the association between the frequency of nocturnal oxygen desaturation, estimated by a pulse oximeter, and the prevalence of atrial fibrillation. We found a significant association between the severity of sleep-disordered breathing (SDB) and the prevalence of atrial fibrillation; the odds ratios (ORs) were 2.47 for those with 5–15 events/h of 3% oxygen desaturation index (ODI) level and 5.66 for those with \geq 15 events/h of 3% ODI level (p for trend = 0.02).

Although the association of SDB with atrial fibrillation has been reported in recent clinical studies, ¹⁻⁴ no population-based epidemiological study has examined this relationship. We investigated the association between the frequency of nocturnal oxygen desaturation and the prevalence of atrial fibrillation among community-based subjects.

METHODS

The subjects were 1763 Japanese men aged 40–74 years who lived in three Japanese communities. They participated in the 2000–2004 annual cardiovascular risk surveys and were recruited for the present sleep study, at a total recruitment rate of 84%. The sensor of the pulse oximeter (PULSOX-3Si, Minolta, Japan) was attached to the index finger during allnight sleep at home. We used a sleep log to exclude waking time from the analysis to minimise potential overestimation of sleep duration. Data from people with total recording time <4 h, or with artefacts likely due to frequent body movement, inadequate fitting of the probe or excessive pulse pressure (n = 100) were excluded, and data from 1663 men were used for the analyses.

A 3% ODI was used to define SDB: 5–<15 events/h was considered mild SDB and ≥15 events/h was considered moderate to severe SDB.⁵ Standard 12-lead electrocardiograms were recorded in the supine position. Each record was coded independently using the Minnesota Code by two trained cardiovascular doctors. The criterion for atrial fibrillation was Minnesota Code 8-3.

To compare the mean values of selected cardiovascular characteristics, we used analysis of covariance with age as covariate. The logistic regression analysis was used to obtain the ORs of the prevalence of atrial fibrillation according to categories of 3% ODI levels (table 1) after adjustment for age, body mass index, alcohol intake, smoking status (never, former smoker, currently 1–19 cigarettes/day and currently ≥20 cigarettes/day), systolic blood pressure and the use of anti-hypertensive drugs. The study was approved by the medical ethics committee of the University of Tsukuba, Tsukuba, Japan, and written informed consent was obtained from all participants.

RESULTS

The mean age, body mass index, alcohol intake, blood pressure, prevalence of use of anti-hypertensive drugs and

Table 1 Age-adjusted means (SEM), prevalence of selected cardiovascular risk characteristics and multivariate-adjusted prevalence of atrial fibrillation in electrocardiogram according to 3% oxygen desaturation index (ODI) among 1663 men

3% ODI		p for	
0–4	5–14	15+	trend
975	534	154	
		61.1 (0.7)	< 0.001
	25.0 (0.1)	26.7 (0.2)	< 0.001
e,22.2 (0.8)	25.2 (1.0)	26.1 (1.9)	0.014
44	38	32	0.001
130.8 (0.5)	133.8 (0.7)	137.6 (1.3)	< 0.001
80.7 (0.3)	82.9 (0.4)	84.9 (0.8)	< 0.001
e18	25	35	< 0.001
42	51	62	< 0.001
7	11	8	
0.8	1.9	4.9	< 0.001
1	2.47 (0.91,	5.66 (1.75,	0.020
	6.69)	18.34)	
	0-4 975 58.7 (0.3) 23.1 (0.1) 2,22.2 (0.8) 44 130.8 (0.5) 80.7 (0.3) e18 42 7 0.8	0-4 5-14 975 534 58.7 (0.3) 59.8 (0.4) 23.1 (0.1) 25.0 (0.1) 2,22.2 (0.8) 25.2 (1.0) 44 38 130.8 (0.5) 133.8 (0.7) 80.7 (0.3) 82.9 (0.4) e18 25 42 51 7 11 0.8 1.9 1 2.47 (0.91,	0-4 5-14 15+ 975 534 154 58.7 (0.3) 59.8 (0.4) 61.1 (0.7) 23.1 (0.1) 25.0 (0.1) 26.7 (0.2) 2,22.2 (0.8) 25.2 (1.0) 26.1 (1.9) 44 38 32 130.8 (0.5) 133.8 (0.7) 137.6 (1.3) 80.7 (0.3) 82.9 (0.4) 84.9 (0.8) e18 25 35 42 51 62 7 11 8 0.8 1.9 4.9 1 2.47 (0.91, 5.66 (1.75,

BMI, body mass index; OR, odds ratio.

*Hypertension was defined as systolic blood pressure \$140 mm Hg, diastolic blood pressure \$90 mm Hg or use of antihypertensive drugs. †Adjusted for age, BMI, alcohol intake, smoking status, systolic blood pressure and use of antihypertensive drugs.

hypertension correlated positively with the 3% ODI level. After adjustment for these covariates, the ORs for atrial fibrillation were 2.47 (95% confidence interval (CI), 0.91 to 6.69) for those with 5–<15 of 3% ODI level (event/h) and 5.66 (95% CI 1.75 to 18.34) for those with \geq 15 of 3% ODI level (p for trend = 0.02).

DISCUSSION

We found a significant positive association between the severity of SDB and the prevalence of atrial fibrillation among community-dwelling Japanese men, independent of cardio-vascular risk factors. The exact mechanisms for the association between SDB and atrial fibrillation are not clear but include chronic intermittent hypoxaemia, hypercapnia, intrathoracic pressure swing and activation of sympathetic nervous function, resulting in cardiac electric instability, distortion of cardiac configuration and hypertension. ¹⁻⁴ SDB is treatable through weight reduction and continuous positive airway pressure. ^{1 2} A recent clinical case report showed that the onset of atrial fibrillation was preceded by a long apnoeic event, and spontaneous reversal to sinus

Abbreviations: ODI, oxygen desaturation index; SDB, sleep-disordered breathing

Oxygen desaturation 1855

rhythm occurred after a period without apnoeas.3 On the other hand, atrial fibrillation usually reduces cardiac output and the reduced cardiac output may lead to central apnoea during sleep, mainly owing to the chemoreflex enhancement and prolonged lag to ventilatory response. 5 We should further clarify whether subjects with atrial fibrillation have a central apnoea pattern by polysomnography.

Atrial fibrillation is a common arrhythmia with associated complications of stroke and other adverse outcomes. Furthermore, atrial fibrillation is an extremely costly public health problem; the direct cost of atrial fibrillation was estimated as 459 million in the year 2000, equivalent to 0.97% of the total National Health Service expenditure.6 This study suggests that the detection of SDB and its successful treatment could be one of the public health approaches to reduce the risk and the cost of atrial fibrillation.

ACKNOWLEDGEMENTS

We thank Ms Minako Tabata and Yukiko Ichikawa for their cooperation.

Authors' affiliations

T Tanigawa, K Yamagishi, S Sakurai, I Muraki, H Noda, Department of Public Health Medicine, Doctoral Program in Social and Environmental Medicine, Graduate School of Comprehensive Human Sciences, University of Tsukuba, Tsukuba, Ibaraki, Japan

T Shimamoto, Osaka Medical Center for Health Science and Promotion, Higashinari-ku, Osaka, Japan

H Iso, Public Health, Department of Social and Environmental Medicine, Graduate School of Medicine, Osaka University, Suita-shi, Osaka,

Funding: This study was supported partly by the Japanese Ministry of Education, Culture, Sports, Science and Technology (grant-in-aid for research B: 14370132), the Health and Labour Sciences Research Grant (Clinical Research for Evidence Based Medicine) and a research grant from FULHAP, Japan. The funding sources had no role in the study design, data collection, data analysis, data interpretation or writing of

Competing interests: None declared.

TT, TS and HI designed the study. TT, KY, SS, HN and IM collected and analysed the data. IM and TT carried out statistical analysis of the data. HI, TT and TS coordinated the study. The manuscript was prepared mainly by TT and HI, with contributions from SS, KY and TS.

Correspondence to: T Tanigawa, Department of Public Health Medicine, Doctoral Program in Social and Environmental Medicine, Graduate School of Comprehensive Human Sciences, University of Tsukuba, 1-1-1, Tennodai, Tsukuba 305-8575, Japan; Tt9178@aol.com

Accepted 9 May 2006

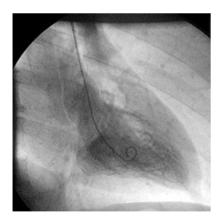
REFERENCES

- Kanagala R, Murali NS, Friedman PA, et al. Obstructive sleep apnea and the recurrence of atrial fibrillation. Circulation 2003;107:2589-94
- Gami AS, Pressman G, Caples SM, et al. Association of atrial fibrillation and obstructive sleep apnea. Circulation 2004;110:364-7.
- Schulz R, Eisele HJ, Seeger W. Nocturnal atrial fibrillation in a patient with
- obstructive sleep apnoed. *Thorax* 2005;**60**:174. **Tanigawa T**, Tachibana N, Yamagishi K, *et al.* Usual alcohol consumption and
- arterial oxygen desaturation during sleep. JAMA 2004;292:923–5.
 Francis DP, Willson K, Davies LC, et al. Quantitative general theory for periodic breathing in chronic heart failure and its clinical implications. Circulation 2000;102:2214–21.
- Stewart S, Murphy N, Walker A, et al. Cost of an emerging epidemic: an economic analysis of atrial fibrillation in the UK. Heart 2004;90:286–92.

IMAGES IN CARDIOLOGY.....

doi: 10.1136/hrt.2006.090183

Unusual diagnosis of ascending aorta dissection with left ventricular angiogram



Left ventricular angiogram after ejection of the contrast product. Note the intimal flap in the ascending aorta between the true and the false lumen. See video 1 to observe retrograde opacification of the false lumen.

49-year-old patient with no medical history was referred to the cardiology department for an acute coronary syndrome. The patient presented with an acute, oppressive chest pain associated with nausea. Physical examination showed symmetric arm blood pressure, all peripheral pulses were present, and there was no evidence of cardiac murmur or abdominal signs. The ECG showed an ST segment elevation on the anterior leads. A coronary angiogram was performed showing normal coronary arteries. The left ventricular angiogram showed an acute dissection of the ascending aorta (see panel and video 1; to view video footage visit the Heart website—http://www.heartjnl.com/supplemental). The delayed retrograde opacification of the false lumen toward the aortic root suggested a site of origin close to the aortic arch. The patient was prepared for emergency cardiac surgery. In the operating room, the patient's pressure collapsed while transoesophageal echocardiography showed a total obliteration of the ascending aorta by the false lumen (videos 2 and 3) resulting in cardiac arrest. Extracorporeal circulation was quickly instituted between the right femoral artery and the right atrium. Surgical treatment consisted of the resuspension of the native aortic valve and sus coronary replacement of the ascending aorta (26 mm Dacron tubular graft). The intimal tear was localised close to the origin of the innominate artery and removed with the ascending aorta. The patient recovered with an uneventful postoperative course. A wide range of clinical manifestations characterise aortic dissection. The diagnosis of ascending aorta dissection should be evocated even in the case of typical acute coronary syndrome.

> T Joudinaud F Baron L Etchegoyen tomtomjoud@hotmail.com

To view video footage visit the Heart website—http:// www.heartinl.com/ supplemental

