



PREVALENCE OF ATRIAL FIBRILLATION AND PRACTICE PATTERNS OF ANTITHROMBOTIC THERAPY IN A POPULATION-BASED COHORT STUDY OF HEMODIALYIS PATIENTS: THE VIENNA INVESTIGATION OF ATRIAL FIBRILLATION AND THROMBOEMBOLISM IN HEMODIALYSIS PATIENTS (VIVALDI)

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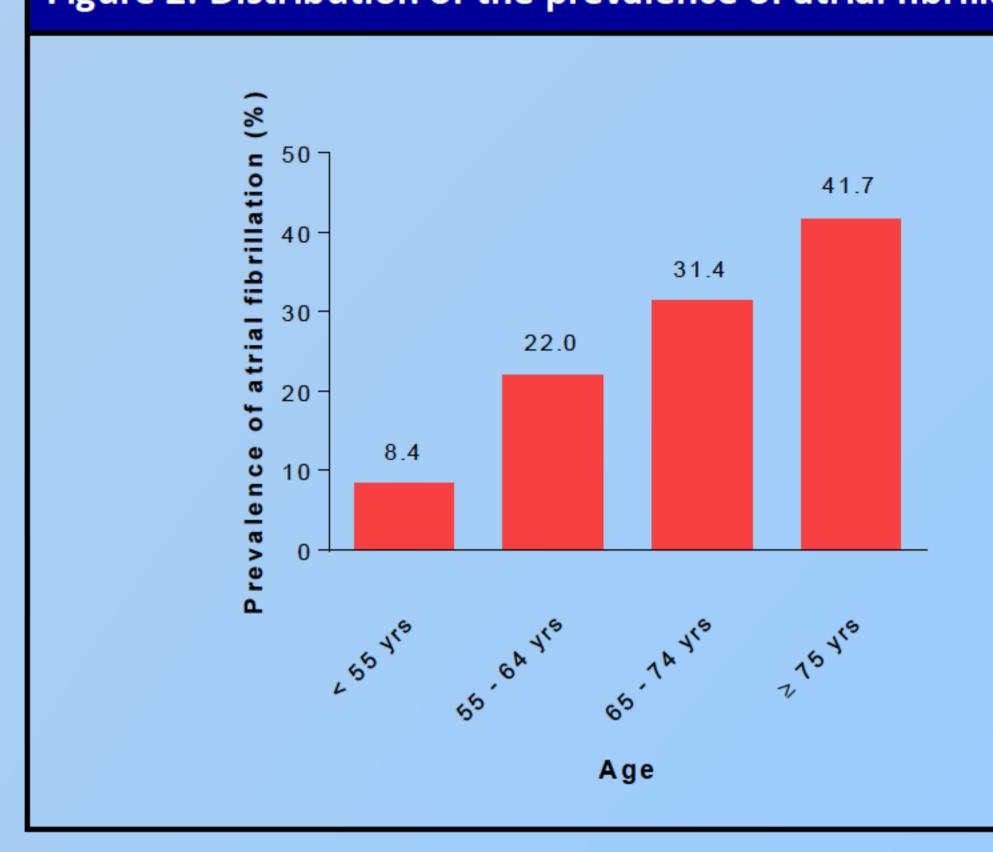
Background:

Atrial fibrillation (AF) is an underestimated comorbidity in hemodialysis (HD) patients that adds significant risk of stroke and thromboembolism to a high burden of disease in end-stage renal disease (ESRD). Data on the prevalence of AF in HD patients derived from patient-level cohort studies are scare. The aim of this study was to investigate the prevalence of AF in a population-based cohort with patient-level data and the distribution of practice patterns of antithrombotic treatment for stroke prevention in AF.

Patients and Methods:

The Vienna InVestigation of AtriaL fibrillation and thromboembolism in patients on hemoDlalysis (VIVALDI) is a population-based cohort study with a cross-sectional baseline data capture phase and an ongoing prospective observational phase, with primary endpoints stroke, thromboembolism, and death. At the seven major hemodialysis centers of Vienna, Austria, 626 patients provided informed consent for participation (figure 1). A structured interview with each individual patient was performed, recent ECGs were viewed and medical histories were verified with the electronic records of the HD centers. A baseline blood sample was taken before hemodialysis and frozen for storage in a state-of-the-art biorepository.

Figure 2: Distribution of the prevalence of atrial fibrillation in the population according to age and HD vintage



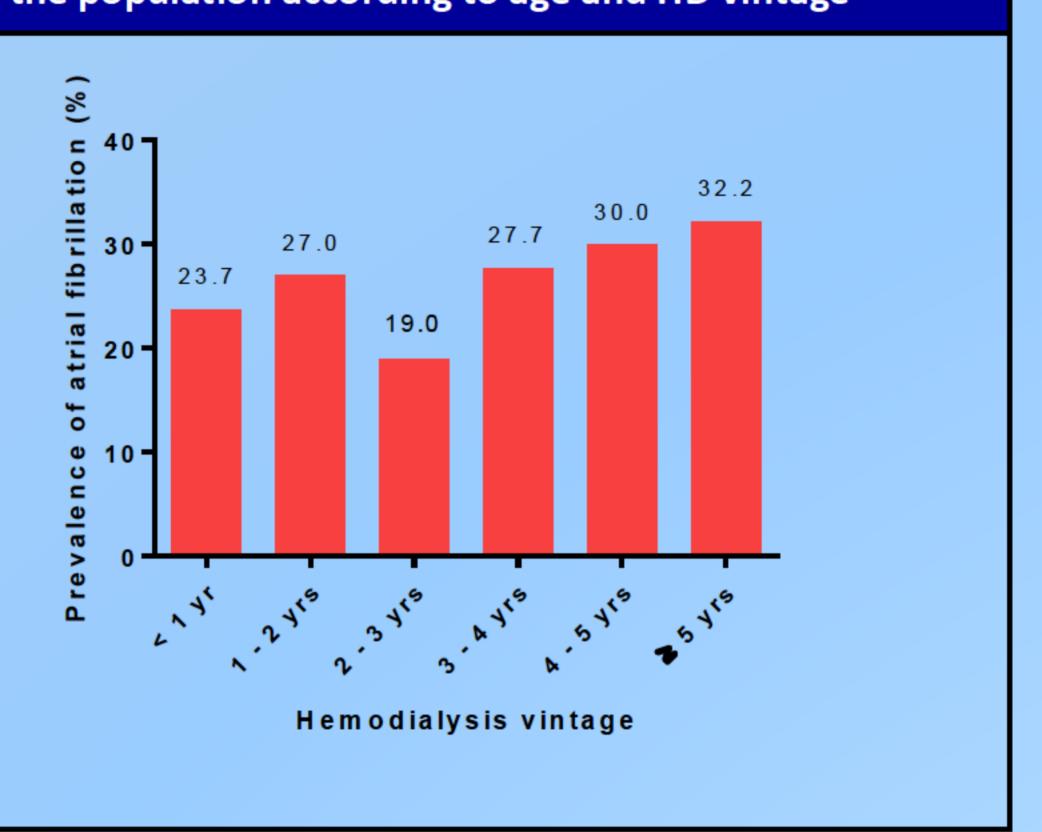


Table 1: Baseline population characteristics Full population Non-AF cohort AF cohort Characteristics p-value* 626 (100) 460 (73.5) 166 (26.5) Patients, n (%) 0.017 397 (63.4) 279 (60.7) Male sex (%) 118 (71.1) 63.5(50 - 73)< 0.001 66 (55 - 75)71.5(64 - 78)Age, median (25th – 75th percentile) BMI, median (25th – 75th 25.7 (22.4 -25.6 (22.2 -25.9 (22.7 – 0.734 29.4) 29.7) percentile) 29.6) Etiology of ESRD, n (%) 160 (25.6) 117 (25.4) 0.906 Diabetic NP 43 (25.9) Vascular NP 121 (19.3) 82 (17.8) 39 (23.5) 0.113 0.203 Glomerular nephritis 82 (13.1) 65 (14.1) 17 (10.2) Atrophic NP 0.781 57 (9.1) 41 (8.9) 16 (9.6) Cystic non-hereditary NP 36 (5.8) 26 (5.7) 0.860 10 (6.0) 0.179 Hereditary NP 31 (5.0) 26 (5.7) 5 (3.0) 0.016 20 (3.2) 10 (2.2) Nephrectomy 10 (6.0) latrogenic/toxic NP 28 (4.5) 19 (4.1) 9 (5.4) 0.491 0.067 Other causes 91 (14.5) 74 (16.1) 17 (10.2) Dialysis history, n (%) History of renal 90 (14.4) 64 (13.9) 0.582 26 (15.7) transplantation 0.606 Previous peritoneal dialysis 46 (7.3) 35 (7.6) 11 (6.6) Current vascular access 329 (52.6) 244 (53.0) 0.684 AV fistula 85 (51.2) 0.246 AV graft 72 (11.5) 57 (12.4) 15 (9.0) 221 (35.3) 0.307 157 (34.1) Central venous catheter 64 (38.6) 0.286 others 4 (0.7) 2 (0.4) 2 (1.2) Dialysis parameters, median (25th – 75th percentile) 0.295 Remaining diuresis, ml 500 (0 – 1000) 500 (0 -1000) 350 (0 – 1000) 1.9(1.2 - 2.6)Inter-dialytic weight gain, kg 2.0 (1.1 – 3.0) 2.1(1.1 - 3.1)0.135 Hemodialysis vintage, years 2.7 (1.0 – 5.0) 2.5(1.0 - 5.0)3.0(1.1-6.0)0.084 Comorbidities, n (%) 82 (17.8) History of stroke or TIA 127 (20.3) 45 (27.1) 0.011 0.048 History of myocardial 105 (16.8) 69 (15.0) 36 (21.7) infarction < 0.001 Coronary heart disease 233 (37.2) 150 (32.6) 83 (50.0) Artificial heart valve 43 (6.9) 22 (4.8) 21 (12.7) 0.001 0.007 History of VTE 61 (9.7) 36 (7.8) 25 (15.1) 0.075 44 (7.0) Deep vein thrombosis 27 (5.9) 17 (10.2) 0.037 Pulmonary embolism 32 (5.1) 18 (3.9) 14 (8.4) Peripheral artery disease 197 (31.5) 139 (30.2) 58 (34.9) 0.261 0.244 Diabetes 237 (37.9) 168 (36.5) 69 (41.6) 576 (92.0) 423 (92.0) 153 (92.2) 0.931 Hypertension < 0.001 Congestive heart failure 184 (29.4) 114 (24.8) 70 (42.2) < 0.001 Cancer history or active 152 (24.3) 96 (20.9) 56 (33.7) Smokers 306 (48.9) 0.351 230 (50.0) 76 (45.8) Current antithrombotic therapy, n (%) LMWH s.c. on non-HD days 59 (9.4) < 0.001 26 (5.7) 33 (19.9) Vitamin K antagonist 27 (5.9) <0.001 77 (12.3) 50 (30.1) Antiplatelet agent 345 (55.1) 0.930 254 (55.2) 91 (54.8)

Results:

The overall prevalence of AF was 26.5% (71.1% male, median age 72 years, 64 - 78) with a median HD vintage of 3 years (table 1). The prevalence of AF increased with the age of study participants (figure 2) and with increasing time on HD treatment, denoted HD vintage. The diagnosis of AF was statistically associated with increased age (odds ratio [OR] 1.05 for every year, 95% confidence interval 1.03 – 1.06), male sex (1.92, 1.18 - 3.14), history of coronary heart disease (2.03, 1.31 – 3.14), presence of artificial heart valves (2.24, 1.02 - 4.89), history of venous thromboembolism (1.94, 1.00 - 3.75), congestive heart failure (1.59, 0.99 - 2.53) and history or active cancer (1.65, 1.02 - 2.68) in univariate logistic regression (table 2). In a multivariable logistic regression model, male sex, age and HD vintage were independently associated with AF (table 2). In 60% of AF patients, AF was diagnosed close to or after ESRD and in 40% it had been present before diagnosis of ESRD. Further baseline population characteristics and their respective associations with AF are shown in table 1.

Antithrombotic treatment was applied in 84.4% of AF patients (28.9% with anticoagulant agents, in 33.7% with antiplatelet agents, and in 21.1% with both). Among the anticoagulant agents used, 60.2% were vitamin-K-antagonists, 38.6% were low-molecular-weight heparins on non-HD-days, and 1.2% were fondaparinux. Twenty-six AF patients (15.6%) received no antithrombotic therapy, except during HD sessions.

Hemodialysis population of Patients excluded Vienna, Austria (N=188): $(N \approx 860)$ Refused consent: 75 Language barriers: 54 Patients approached and offered Cognitive impairment: participation Guests at dialysis (N = 814)center: 7 Withdrawn consent: Patients included: (N = 626) 1 patient Other reasons: 34

Figure 1: Flowchart of patient recruitment

12-lead ECG
Medical records search
Blood sampling for biorepository

AF
No AF
N = 166
N = 460

Baseline Data Capture:

Structured interview

26.5%

WILHELMINENSPITAL

DIE MENSCHIEßE GFÖGE

Aug DONAUSPITAL

Dialyse mit
Perspektive

Dialyse mit
Perspektive

Dialyse mit
Perspektive

Auguste der Stadische Größe

Auguste der Stadische Größe

Wener Stadische Größe

Auguste der Stadische Größe

Donaumer

Wener Stadische Größe

Donaumer

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Wener Stadische Größe

Auguste der Größe

Donaumer

Donaumer

Wener Stadische Größe

Auguste Größe

Donaumer

Donaumer

Wener Dialyse mit
Perspektive

Auguste Größe

Donaumer

Donaumer

Donaumer

Donaumer

Neuer Stadische Größe

Auguste Größe

Donaumer

Donaumer

Donaumer

Donaumer

Donaumer

REDLING

REDLING

RAISER FRANZ JOSEF SPITAL

DONAUMER DONAUM KO

73.5%

Table 2: Logistic regression model of factors associated with atrial fibrillation (N=559)

Risk factors	Univariable OR	р	Multivariable OR	р
	(95%CI)		(95%CI)	
Male sex	1.92 (1.18 - 3.14)	0.009	2.18 (1.28 - 3.70)	0.004
Age, per year	1.05 (1.03 - 1.06)	<0.001	1.05 (1.03 - 1.07)	<0.001
HD vintage, per year	1.06 (1.02 - 1.11)	0.003	1.08 (1.03 - 1.13)	0.001
Stroke/TIA	1.56 (0.93 - 2.60)	0.090	1.14 (0.66 - 1.98)	0.631
Coronary heart disease	2.03 (1.31 - 3.14)	0.002	1.41 (0.87 - 2.27)	0.160
Artificial heart valve	2.24 (1.02 - 4.89)	0.043	1.44 (0.62 - 3.34)	0.400
VTE	1.94 (1.00 - 3.75)	0.049	1.89 (0.93 - 3.85)	0.081
Congestive heart failure	1.59 (0.99 - 2.53)	0.052	1.24 (0.75 - 2.03)	0.403
Cancer history/active	1.65 (1.02 - 2.68)	0.043	1.21 (0.72 - 2.03)	0.481

Conclusions:

The prevalence of AF is high amongst HD patients and seems to increase with HD vintage. AF is associated with patient age, sex and distinct medical comorbidities. Practice patterns of antithrombotic treatment indicate the expected lack of consensus for stroke prevention in AF for HD patients. Evidence-based and HD-specific risk evaluation strategies are crucial for guiding initiation and duration of anticoagulation treatment in this high-risk population.

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