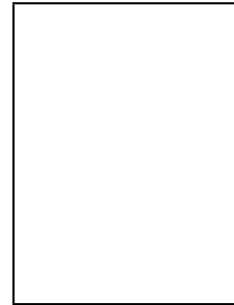


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Non-vitamin K versus vitamin K antagonist oral anticoagulants in surgical mitral valve repair or bioprosthetic valve replacement in the first three months after surgery

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Abstract

Introduction and Objectives: Oral anticoagulation (OAC) with non-vitamin K antagonist oral anticoagulants (NOACs) after surgical mitral valve repair (MVR) or bioprosthetic valve replacement (BVR) in mitral position remains a controversial topic among the cardiovascular community, in particular in the early postoperative period. This study aimed to evaluate the efficacy and safety of NOACs in the first three months after MVR or mitral BVR compared to vitamin K antagonists (VKAs).

Methods: This was a single-center retrospective study with prospectively collected peri-intervention outcomes between 2020 and 2021. Records were retrieved and all participants were contacted by telephone. Patients were divided into groups according to OAC strategy. The primary outcome was a composite of death, rehospitalization, myocardial infarction, stroke or transient ischemic attack, systemic embolism, mitral thrombosis, or bleeding during the first three months after surgery.

Results: A total of 148 patients were enrolled, with a mean age of 65.5 ± 12.2 years, 56.8% male. On discharge, 98 (66.2%) patients were on VKAs and 50 (33.8%) were on DOACs for at least three months. The primary outcome occurred in 22 (22.4%) patients in the VKA group and in three (6%) in the NOAC group ($p=0.012$), mainly driven by more bleeding events in the former. Independent predictors of the primary outcome were smoking ($p=0.028$) and OAC with VKAs at discharge, the latter predicting three times more events ($p=0.046$, OR 3.72, 95% CI 1.02-13.5).

Conclusions: NOACs were associated with fewer events, supporting their efficacy and safety during the first three months after surgical MVR or mitral BVR.

KEYWORDS

Non-vitamin K antagonist oral anticoagulants; Mitral valve repair; Mitral bioprosthetic valve replacement; First three months after surgery

Anticoagulantes orais não antagonistas da vitamina K versus antagonistas da vitamina K nos primeiros três meses após reparação cirúrgica da válvula mitral ou substituição cirúrgica por prótese valvular biológica em posição mitral

Resumo

Introdução e objetivos: A anticoagulação oral (OAC) com anticoagulantes orais não antagonistas da vitamina K (NOACs) após reparação cirúrgica da valva mitral (MVR) ou substituição cirúrgica por prótese valvular biológica (BVR) em posição mitral permanece um tópico controverso entre a comunidade cardiovascular, em particular no período pós-operatório inicial. O objetivo do estudo foi avaliar a eficácia e segurança dos NOACs nos primeiros três meses após MVR ou BVR em posição mitral, comparando esta estratégia com antagonistas da vitamina K (AVKs).

Métodos: Análise retrospectiva de um único centro com resultados peri-intervenção colhidos prospectivamente entre 2020/2021. Os registros foram analisados e todos os participantes foram contatados por telefone. Os pacientes foram divididos em grupos de acordo com a estratégia OAC. O outcome primário composto foi definido como morte, re-hospitalização, enfarte do miocárdio, acidente vascular cerebral ou ataque isquêmico transitório, embolia sistêmica, trombose mitral ou hemorragia durante os primeiros três meses após a cirurgia.

Resultados: Foram incluídos 148 doentes, com uma idade média de $65,5 \pm 12,2$ anos e 56,8% eram do sexo masculino. À data da alta, 98 (66,2%) pacientes foram medicados com AVKs e 50 (33,8%) com NOACs, mantidos pelo menos durante três meses. O outcome primário ocorreu em 22 (22,4%) pacientes no grupo AVKs e em 3 (6%) no grupo NOACs ($p=0,012$), principalmente sobre a influência de um maior número de eventos hemorrágicos no primeiro. Os preditores independentes do outcome primário foram tabagismo ($p=0,028$) e OAC com AVKs à data da alta, este último prevendo três vezes mais eventos ($p=0,046$, OR 3,72, IC 95% 1,02 a 13,5).

Conclusões: Os NOACs foram associados a menos eventos, nomeadamente hemorrágicos, apoiando a sua eficácia e segurança durante os primeiros três meses em doentes após MVR ou BVR em posição mitral.

PALAVRAS-CHAVE

Anticoagulantes orais não antagonistas da vitamina K; Reparação cirúrgica da valva mitral; Implantação cirúrgica de prótese valvular em posição mitral; Primeiros três meses após a cirurgia.

Introduction

Valvular heart disease affects more than 100 million people worldwide and an estimated 300 000 prosthetic heart valves are implanted every year. This rate is increasing, mainly driven by intrinsic valvular degeneration in aging populations.¹

The use of bioprosthetic valve replacement (BVR) has also increased in the last 20 years compared to mechanical heart valves. This shift is difficult to explain, but may be related to the drawbacks associated with long-term oral anticoagulation (OAC) in younger patients as well as the higher burden of side effects, particularly in the elderly.² Even with appropriate OAC therapy after valve implantation, there is a higher lifelong risk of thromboembolic events (estimated at 1-4%) and bleeding (2-9%) compared to the general population.^{3,4}

Non-vitamin K antagonist oral anticoagulants (NOACs) have been shown to be safe and at least as effective as vitamin K antagonists (VKAs) in patients with atrial fibrillation (AF) and without moderate to severe mitral stenosis or rheumatic valvular disease (RVD).⁵ However, patients with mechanical heart valves should be anticoagulated with VKAs, given the harmful effects of dabigatran noted in the REALIGN phase II trial, and more recently, the premature termination of the PROACT Xa trial of apixaban versus warfarin.⁵⁻⁷

However, the ideal OAC strategy in patients with mitral BVR or mitral valve repair (MVR) remains a matter of debate in the cardiovascular community, especially regarding the first three months after surgery. The American College of Cardiology/American Heart Association (ACC/AHA) guidelines recommend initiating OAC with VKAs within three months of index valvular surgery regardless of rhythm or valve position, and the European Society of Cardiology/European Association for Cardio-Thoracic Surgery (ESC/EACTS) guidelines also recommend VKAs for biological heart valves in mitral position within three months, adding that NOACs may be considered for patients with AF and a bioprosthesis in mitral position (class of recommendation IIb, level of evidence C). These recommendations are mainly based on observational studies.⁸⁻¹²

Nevertheless, off-label use of NOACs within three months of index mitral surgery is growing, supported by small studies and meta-analyses.^{4,13-15}

In the RIVER trial, the subgroup of patients taking rivaroxaban in the first months after index valvular surgery showed fewer primary endpoint events, but only 18.8% of the patients in the main study were randomized within three months.¹⁶ Similarly, the ENAVLE trial showed that edoxaban was non-inferior

to warfarin regarding the primary efficacy endpoint as well as for major bleeding. However, this was a small trial which included patients with bioprosthetic aortic valve replacement (49%), which may limit the applicability of these results.¹⁷

Regarding thromboembolic and bleeding events, NOACs appear to be as safe and as effective as VKAs in patients with surgical mitral BVR or MVR within three months of surgery. However, stronger evidence from observational studies and larger randomized trials are needed to further sustain these findings.

Objectives

This study aimed to evaluate the efficacy and safety of NOACs in the first three months after MVR or mitral BVR compared to VKAs.

Methods

Ethical statement

The study abided by the principles stated in the 1975 Helsinki Declaration and was approved by the institutional ethics committee of the hospital center in which the study took place.

Patient selection

This was a single-center retrospective cohort study, with prospectively collected peri-intervention outcomes between 2020 and 2021, which included patients undergoing either surgical MVR or mitral BVR.

Patients were eligible for the study if they underwent MVR or mitral BVR. Patients who underwent mechanical prosthetic valve implantation or concomitant ascending aorta replacement or other valvular procedure, or who died during the index hospitalization, were excluded. Patients treated by other antithrombotic strategies such as concomitant use of antiplatelets or heparin were also excluded.

Patients were then divided into two groups according to OAC strategy (NOACs or VKAs). In patients receiving VKAs on discharge, the recommended target international normalized ratio (INR) was between 2 and 3. They were entered into a follow-up period of at least three months, with monitoring at specific time points (either at the hospital or with their primary care provider). Baseline patient demographic data, cardiovascular risk factors, and clinical, laboratory, echocardiographic and surgical data were recorded. All patients were contacted by telephone. Switching OAC therapy only occurred during hospitalization, and it was maintained in the postoperative period.

Outcomes

The primary outcome (net clinical benefits) was a composite of death, rehospitalization, myocardial infarction, stroke or transient ischemic attack, systemic embolism, mitral thrombosis, or bleeding during the first three months after surgery. Bleeding events were defined according to the Bleeding Academic Research Consortium (BARC) scale.¹⁸ A separate analysis aimed to assess bleeding (on the BARC scale) and stroke events in the AF population during the first three months after surgery.

Statistical analysis

Continuous variables were presented as mean \pm standard deviation or median and interquartile range. These were compared between groups using the independent samples t test or the Mann-Whitney U test, based on their distribution. Categorical variables were presented as frequencies and percentages and were compared using the chi-square test or Fisher's exact test, as appropriate.

Separate analyses were conducted for net clinical benefits (primary outcome). Independent predictors of the primary outcome were assessed by multivariate logistic regression analyses using the forward stepwise method. The Hosmer-Lemeshow test was used to calibrate the regression model. The effects of the variables were assessed by estimating odds ratios (OR) and 95% confidence intervals (CI). The variables entered into the model were gender, age, smoking status, history of type 2 diabetes, and VKAs at discharge.

In the AF subpopulation, a separate analysis for bleeding events (on the BARC scale) was also conducted. The variables entered into the model were gender, age, anticoagulation, and switching OAC therapy at discharge. Statistical differences between the presence of AF, type of surgery, and presence of RVD were assessed by analysis of variance (ANOVA). A p-value for interaction of <0.05 was taken to indicate statistical significance.

Results

Baseline patient data

A total of 170 patients were initially identified, of whom 22 were excluded due to death during hospitalization (n=9) or mechanical mitral valve surgery (incorrectly classified) (n=13) (Figure 1). A total of 148 patients were therefore eligible and were contacted.

Participants' mean age was 65.5 ± 12.2 years and 56.8% of the sample were male. In terms of comorbidities, 73% were hypertensive, 14.9% were diabetic, 19.6% were obese, 9.5% had chronic kidney disease and 45.3% had heart failure. Clinically, AF was previously diagnosed in 60 (40.5%) patients.

Fifty-nine (39.9%) were on OAC therapy before surgery: 22 (14.9%) with VKAs and 37 (25%) with DOACs. RVD was present in 12.2%. From a surgical standpoint, 76 (51.4%) patients underwent surgical mitral BVR and 72 (48.6%) MVR. On discharge, 98 (66.2%) were on VKAs and 50 (33.8%) were on NOACs maintained for at least three months (Table 1).

Table 1 Patient demographic, clinical, and surgical characteristics compared by anticoagulation strategy at discharge.

	Anticoagulation			p
	VKAs (n=98, 66.2%)	NOACs (n=50, 33.8%)	Total (n=148)	
Gender, n (%)				0.204
Male	52 (53.1)	32 (64.0)	84 (56.8)	
Female	46.0 (46.9)	18.0 (36)	64.0 (43.2)	
Age, years, mean \pm SD	66.2 \pm 11.7	64.5 \pm 13.0	65.5 \pm 12.2	0.413
Weight, kg, mean \pm SD	73.0 \pm 14.2	72.6 \pm 12.7	73.0 \pm 13.7	0.879
Height, cm, mean \pm SD	166 \pm 9.0	167 \pm 10.0	166 \pm 10.0	0.416
BMI, kg/m ² , mean \pm SD	26.2 \pm 5.20	25.8 \pm 4.10	26.1 \pm 4.80	0.666
Hypertension, n (%)	71 (72.4)	37 (74.0)	108 (73.0)	0.841
Type 2 diabetes, n (%)	17 (17.3)	50 (10.0)	22 (14.9)	0.235
Dyslipidemia, n (%)	41 (41.8)	16 (32.0)	57 (38.5)	0.245
Obesity, n (%)	21 (21.4)	80 (16.0)	29 (19.6)	0.431
Previous stroke, n (%)	0 (0.0)	1 (2.0)	1 (0.70)	0.160
Smoking, n (%)	18 (18.4)	40 (8.0)	22 (14.9)	0.094
Chronic renal disease, n (%)	9 (9.20)	5 (10.0)	14 (9.50)	0.872
Atrial fibrillation, n (%)	39 (39.8)	21 (42.0)	60 (40.5)	0.796
Heart failure, n (%)	43 (43.9)	24 (48.0)	67 (45.3)	0.634
VTE, n (%)	1 (1.0)	1 (2.0)	2 (1.40)	0.625
Previous anticoagulation, n (%)	38 (38.8)	21 (42.0)	59 (39.9)	0.705
Previous anticoagulant class, n (%)				
VKAs	21 (55.3)	1 (4.80)	22 (37.3)	
NOACs	17 (44.7)	20 (95.2)	37 (62.7)	<0.001
Total	38 (38.8)	21 (42.0)	59 (39.9)	
Switch (NOACs to VKAs) at discharge, n (%)	17 (17.3)	1 (2.0)	18 (12.2)	0.007
Previous drug, n (%)				
Apixaban	5 (13.2)	3 (14.3)	8 (13.6)	
Dabigatran	3 (7.90)	4 (19.0)	7 (11.9)	
Edoxaban	4 (10.5)	8 (38.1)	12.0 (20.3)	
Rivaroxaban	21 (55.3)	1 (4.80)	22 (37.3)	
Warfarin	38 (38.8)	21 (42.0)	59 (39.9)	0.002
Total	58 (59.2)	18 (36.0)	76 (51.4)	
Type of surgery, n (%)				0.008
Mitral bioprothesis				
MVR	40 (40.8)	32 (64.0)	72 (48.6)	
LAA closure, n (%)	5 (5.10)	3 (6.0)	8 (5.40)	0.819

RVD, n (%)	15 (15.3)	3 (6.0)	18 (12.2)	0.101
LVEF, mean \pm SD	54.6 \pm 8.98	56.2 \pm 8.43	55.4 \pm 8.76	0.222

BMI: body mass index; LAA: left atrial appendage; LVEF: left ventricular ejection fraction; MVR: mitral valve repair; NOACs: non-vitamin K antagonist oral anticoagulants; RVD: rheumatic valve disease; SD: standard deviation; VKAs: vitamin k antagonists; VTE: venous thromboembolism.

Outcomes

The primary composite outcome occurred in 22 patients (22.4%) in the VKA group and in three patients (6%) in the NOAC group ($p=0.012$). Analysis of the individual components of the primary outcome shows that there were no differences in mortality ($p=0.104$), rehospitalization ($p=0.987$), myocardial infarction (no events in either group), stroke or transient ischemic attack ($p=0.507$), systemic embolism (no events in either group) or mitral thrombosis (no events in either group). The main difference in the primary outcome was driven by more bleeding events (BARC 1 bleeding) in the VKA group ($n=19$, 19.4% vs. $n=2$, 4% in the NOAC group) ($p=0.011$) (Table 2). No BARC >1 events were recorded during the follow-up period.

Table 2 Primary outcome and its individual components compared by anticoagulation strategy at discharge.

	Anticoagulation			p
	VKAs (n=98, 66.2%)	NOACs (n=50, 33.8%)	Total (n=148)	
Primary outcome, n (%)	22 (22.4)	30 (60)	25 (16.9)	0.012
Individual components, n (%)				
Death	50 (5.10)	0 (0)	50 (3.40)	0.104
Rehospitalization	20 (20)	10 (20)	30 (20)	0.987
Myocardial infarction	0 (0)	0 (0)	0 (0)	^a
Stroke or TIA	40 (4.10)	10 (20)	50 (3.40)	0.507
Systemic embolism	0 (0)	0 (0)	0 (0)	^a
Mitral thrombus	0 (0)	0 (0)	0 (0)	^a
BARC 1 bleeding	19 (19.4)	20 (40)	21 (14.2)	0.011

BARC: Bleeding Academic Research Consortium criteria; NOACs: non-vitamin K antagonist oral anticoagulants; TIA: transient ischemic attack; VKAs: vitamin K antagonists.

^a No events in either group.

In a subgroup analysis, no statistically significant differences were found for the presence of AF (p for interaction=0.823), type of surgery (p for interaction=0.954) or presence of RVD (p for interaction=0.171) with regard to the primary outcome.

Independent predictors of the primary outcome

In logistic regression analysis, independent predictors of the primary outcome were smoking ($p=0.028$) and OAC with VKAs on discharge ($p=0.046$); the latter was predicted to increase the chance of primary

outcome events threefold ($p=0.046$, OR 3.72, 95% CI 1.02-13.5) (Figure 2).

Outcomes in atrial fibrillation patients

A total of 60 (40.5%) patients with AF were identified, of whom 59 (98.3%) were on OAC therapy before surgery, 22 (37.3%) with VKAs and 37 (62.7%) with NOACs. At discharge, 39 (65%) were on VKAs and 21 (35%) were on NOACs maintained for at least three months. Switching OAC therapy on discharge occurred more frequently in the VKA group: of the 39 patients on VKAs at discharge, 17 (43.6%) patients were previously treated with NOACs. By contrast, of the 21 patients discharged with NOACs, only one (4.8%) was previously on VKAs ($p=0.002$) (Table 3). RVD was present in 16 (26.7%) patients, of whom 14 (87.5%) were discharged on VKAs ($p=0.028$).

Table 3 Anticoagulation regimes of atrial fibrillation patients undergoing surgical mitral valve repair or mitral bioprosthetic valve replacement, compared by anticoagulation strategy at discharge.

	VKAs (n=39, 65%)	Anticoagulation		p
		NOACs (n=21, 35%)	Total (n=60)	
Previous anticoagulation, n (%)	38 (97.4)	21 (100)	59 (98.3)	0.459
Previous drug class, n (%)	21 (55.3)	10 (4.80)	22 (37.3)	
VKAs				
NOACs	17 (44.7)	20 (95.2)	37 (62.7)	
Total	38 (97.4)	21 (100)	59 (98.3)	<0.001
Switch (NOACs to VKAs) at discharge, n (%)	17 (43.6)	10 (4.80)	18 (30)	0.002

NOACs: non-vitamin K antagonist oral anticoagulants; VKAs: vitamin K antagonists.

Bleeding events classified as BARC 1 occurred in the VKA group only (six patients, 15.4%) ($p=0.058$). Ischemic stroke in the three-month postoperative period occurred in two patients (5.1%) on VKAs and in one patient (4.8%) in the NOAC group ($p=0.950$) (Table 4). There was no statistical difference for bleeding (p for interaction=0.119) or stroke events (p for interaction=0.789) regarding the presence of RVD.

Table 4 Bleeding and stroke events in atrial fibrillation patients undergoing surgical mitral valve repair or mitral bioprosthetic valve replacement, compared by anticoagulation strategy at discharge.

	VKAs (n=39, 65%)	Anticoagulation		p
		NOACs (n=21, 35%)	Total (n=60)	
Primary outcome (BARC 1 bleeding), n (%)	6 (15.4)	0 (0)	6 (10.0)	0.058
Secondary outcome (stroke), n (%)	2 (5.1)	1 (4.8)	3 (5.0)	0.950

BARC: Bleeding Academic Research Consortium criteria; NOACs: non-vitamin K antagonist oral anticoagulants; VKAs: vitamin K antagonists.

Previous anticoagulation ($p=0.002$) and switching OAC therapy on discharge from NOACs to VKAs

were associated with bleeding ($p=0.039$), the latter predicting 12 times more events ($p=0.039$, OR 12.4, 95% CI 1.13-134) (Table 5).

Table 5 Multivariate analysis (logistic regression) for bleeding events in atrial fibrillation patients.

	OR (95% CI)	p
Age	0.95 (0.85-1.05)	0.329
Gender	0.89 (0.11-7.00)	0.912
Switch (NOACs to VKAs)	12.4 (1.13-134)	0.039
Anticoagulation	1.00 (0.98-1.02)	1.000

CI: confidence interval; NOACs: non-vitamin K antagonist oral anticoagulants; OR: odds ratio; VKAs: vitamin K antagonists.

Discussion

This study found that treatment with NOACs within three months of bioprosthetic mitral valve implantation or mitral repair was associated with fewer net clinical events, mainly driven by less bleeding. In addition, switching OAC strategy from NOACs to VKAs at discharge in AF patients was associated with more bleeding events. Therefore, these results suggest that NOACs were as safe and as effective as VKAs in this patient cohort.

In the present study, cardiovascular risk factors such as hypertension, dyslipidemia and diabetes resembled those of the groups studied in the RIVER and ENAVLE trials.^{16,17} From a surgical standpoint, 51.4% of patients underwent mitral BVR and 48.6% MVR; AF was previously diagnosed in 40.5% of patients.

At discharge, 98 (66.2%) patients were on VKAs and 50 (33.8%) on NOACs for at least three months. In the subgroup of AF patients with previous indication for OAC, therapy on admission influenced the OAC strategy on discharge, with 21 patients (55.3%) in the VKA group and 20 (95.2%) in the NOAC group discharged under the same therapy prescribed on admission. Switching OAC on discharge was infrequent (18 patients, 30%), albeit more common in the VKA group (43.6% patients switched from NOACs to VKAs on discharge, $p=0.002$).

The primary outcome occurred in 22 (22.4%) patients in the VKA group and in three (6%) in the NOAC group ($p=0.012$), mainly driven by more bleeding events in the former (19.4%, $p=0.011$). No differences were found with regard to death, rehospitalization, myocardial infarction, stroke or transient ischemic attack, systemic embolism or mitral thrombosis. Similarly, no statistically significant difference was found for the presence of AF (p for interaction=0.823).

Bleeding events were classified as 1 on the BARC criteria scale, and OAC with VKAs on discharge

was predicted to generate three times more events. The bleeding rate in the VKA group was similar to the subgroup of patients with bioprosthetic valves included in the trials that compared NOACs to VKAs, although driven by minor bleeding events.^{16,19–22} This was probably related to the short follow-up period (within three months of surgery) in the study, since NOAC-related bleeding events tend to occur later in the course of treatment.²³

The primary efficacy outcome in the ENAVLE trial (death, clinical thromboembolic event, or asymptomatic intracardiac thrombus) favored edoxaban (noninferior and superior) over warfarin, as did safety outcomes (bleeding events) and net clinical outcome, but without statistical difference (noninferior).¹⁷ These results were consistent with those in the present study, with some differences: our study showed NOACs to have comparable efficacy and better safety properties compared to VKAs, whereas ENAVLE showed better efficacy and similar safety in patients treated with NOACs; the lower rate of edoxaban use in the present study does not permit a direct comparison between the studies.

Analysis of the subpopulation of AF patients in the present study showed no differences in bleeding or stroke events in the NOAC group compared to VKA patients, although it is suggestive of fewer bleeding events in the former. In the RIVER trial,¹⁶ the primary outcome (net clinical outcome) was noninferior for rivaroxaban versus warfarin, and bleeding events also showed no differences between groups, although only 18.8% of patients were enrolled and randomized within three months of surgery. These findings were corroborated by the present subanalysis of AF patients.

Transition between different types of oral OAC may represent a period of increased risk of thromboembolism or bleeding, especially when switching from NOACs to VKAs.^{24,25} In the ROCKET-AF trial,²² there was an increase in thrombotic events after discontinuation of rivaroxaban and transition to warfarin at the end of the study. At the end of the ARISTOTLE trial²⁶ (apixaban versus warfarin in AF patients), the blinded study drug was stopped, and open label warfarin was recommended. Patients who switched from apixaban to warfarin had more thrombotic and bleeding events. The higher risk of stroke and bleeding in these patients was probably associated with initiation of VKAs (comparable with the group on stable warfarin) rather than with stopping apixaban.²⁴ Similarly, in the present study, switching OAC therapy on discharge in AF patients from NOACs to VKAs was associated with 12-fold higher odds of more bleeding ($p=0.039$).

While the current 2020 ACC/AHA guidelines provide no recommendation on OAC after MVR, the ESC/EATCS guidelines recommend thromboprophylaxis with VKAs in the first three months, as well as

in patients undergoing mitral BVR.^{8,9} The clinical evidence for antithrombotic therapy after MVR is limited, based mainly on expert opinion, but the use of an annuloplasty ring and the possible occurrence of AF after surgery may justify short-term therapy.²⁷ From a surgical standpoint, the present analysis showed that 64% of those in the NOAC group underwent MVR and 59.2% of the VKA group underwent mitral BVR ($p=0.008$), and the primary outcome showed no statistical difference between the type of surgery (p for interaction= 0.954). NOACs thus appear to be a potential alternative to VKAs.

Although rate and rhythm management of rheumatic AF after valve intervention are suggested in the literature, understanding of the management of antithrombotic therapy after MVR or mitral BVR in RVD patients is limited.²⁸ In this study, RVD was present in 26.7% patients, of whom 87.5% were discharged on VKAs ($p=0.028$). Nevertheless, bleeding (p for interaction= 0.119) and stroke events (p for interaction= 0.789) did not differ depending on the presence of RVD.

Limitations

This was an observational study based on information obtained from medical records and by telephone consultation with patients, and is thus limited in the scope of collectable data. For example, lack of reporting of INR and time in therapeutic range is an acknowledged limitation. The presence of only minor bleeding events (BARC 1) may also limit the significance of the study's conclusions. Finally, the findings from this study, which recruited a small sample from a single center, may not be generalizable to other cardiac centers.

Conclusion

NOACs were associated with fewer events within three months of surgical MVR or mitral BVR, supporting their efficacy and safety compared to VKAs. In addition, recent data from large trials predict a promising role for NOACs used in the immediate postoperative period in these patients.

Conflicts of interest

The authors have no conflicts of interest to declare.

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Ethics in publishing

1. Does your research involve experimentation on animals?:

No

2. Does your study include human subjects?:

Yes

If yes; please provide name of the ethical committee approving these experiments and the registration number. :

Ethical committee of Algarve University and Hospital Center (CHUA)

If yes; please confirm authors compliance with all relevant ethical regulations. :

Yes

If yes; please confirm that written consent has been obtained from all patients. :

Yes

3. Does your study include a clinical trial?:

No

4. Are all data shown in the figures and tables also shown in the text of the Results section and discussed in the Conclusions?:

Yes

References

1. Sun JC, Davidson MJ, Lamy A, et al. Antithrombotic management of patients with prosthetic heart valves: current evidence and future trends. *Lancet*. 2009;374(9689):565–76. Available from: [http://dx.doi.org/10.1016/S0140-6736\(09\)60780-7](http://dx.doi.org/10.1016/S0140-6736(09)60780-7)
2. Brown JM, O'Brien SM, Wu C, et al. Isolated aortic valve replacement in North America comprising 108,687 patients in 10 years: Changes in risks, valve types, and outcomes in the Society of Thoracic Surgeons National Database. *J Thorac Cardiovasc Surg*. 2009;137(1):82–90.
3. Durães AR, de Souza Roriz P, de Almeida Nunes B, et al. Dabigatran Versus Warfarin After Bioprosthesis Valve Replacement for the Management of Atrial Fibrillation Postoperatively: DAWA Pilot Study. *Drugs R D*. 2016;16(2):149–54.
4. Magro PL, Sousa-Uva M. Direct oral anticoagulants and surgical bioprosthetic valves: State of the art. *Rev Port Cardiol*. 2023;42:179–81.
5. Caldeira D, David C, Costa J, et al. Non-Vitamin K antagonist oral anticoagulants in patients with atrial fibrillation and valvular heart disease: Systematic review and meta-analysis. *Eur Hear J - Cardiovasc Pharmacother*. 2018;4(2):111–8.
6. Eikelboom JW, Connolly SJ, Brueckmann M, et al. Dabigatran vs. Warfarin in patients with mechanical heart valves. *N Engl J Med*. 2013;(26).
7. Svensson L, Wang TY, Wen J, et al. PROACT Xa: A Multicenter, Randomized Clinical Trial to Evaluate the Efficacy and Safety of Apixaban vs. Warfarin in Patients with a Mechanical Bileaflet

- Aortic Heart Valve. *Am Assoc Thorac Surg*. 2022;(17):13–4.
8. Otto CM, Nishimura RA, Bonow RO, et al. 2020 ACC/AHA Guideline for the Management of Patients With Valvular Heart Disease: A Report of the American College of Cardiology/American Heart Association Joint Committee on Clinical Practice Guidelines. Vol. 143, *Circulation*. 2021. 72–227 p.
 9. Vahanian A, Beyersdorf F, Praz F, et al. 2021 ESC/EACTS Guidelines for the management of valvular heart disease: Developed by the Task Force for the management of valvular heart disease of the European Society of Cardiology (ESC) and the European Association for Cardio-Thoracic Surgery (EACTS). *Rev Esp Cardiol (Engl Ed)*. 2022;75(6):524.
 10. ElBardissi AW, DiBardino DJ, Chen FY, et al. Is early antithrombotic therapy necessary in patients with bioprosthetic aortic valves in normal sinus rhythm? *J Thorac Cardiovasc Surg*. 2010;139(5):1137–45. Available from: <http://dx.doi.org/10.1016/j.jtcvs.2009.10.064>
 11. Sundt TM, Zehr KJ, Dearani JA, et al. Is early anticoagulation with warfarin necessary after bioprosthetic aortic valve replacement? *J Thorac Cardiovasc Surg*. 2005;129(5):1024–31.
 12. Heras M, Chesebro JH, Fuster V, et al. High risk of thromboemboli early after bioprosthetic cardiac valve replacement. *J Am Coll Cardiol*. 1995;25(5):1111–9.
 13. Magro PL, Sousa-Uva M. Are NOACs as safe and efficient as VKA regarding thromboembolic prophylaxis and major bleeding in patients with surgical bioprosthesis and atrial fibrillation within 3 months of surgery? *Interact Cardiovasc Thorac Surg*. 2022;34(5):739–43.
 14. Eikelboom R, Whitlock RP, Muzaffar R, et al. Direct oral anticoagulants versus vitamin K antagonists in the first 3 months after bioprosthetic valve replacement: a systematic review and meta-analysis. *Eur J Cardio-thoracic Surg*. 2023;63(4):5–8.
 15. Saade W, Peruzzi M, Biondi-Zoccai G, et al. Is it the time for direct oral anticoagulants in bioprosthetic heart valves? *Eur J Cardio-thoracic Surg*. 2023;63(4):90–1.
 16. Guimarães HP, Lopes RD, de Barros e Silva PGM, et al. Rivaroxaban in Patients with Atrial Fibrillation and a Bioprosthetic Mitral Valve. *N Engl J Med*. 2020;383(22):2117–26.
 17. Shim CY, Seo J, Kim YJ, et al. Efficacy and safety of edoxaban in patients early after surgical bioprosthetic valve implantation or valve repair: A randomized clinical trial. *J Thorac Cardiovasc Surg*. 2023;165(1):58-67.e4.
 18. Mehran R, Rao S V., Bhatt DL, et al. Standardized bleeding definitions for cardiovascular clinical trials: A consensus report from the Bleeding Academic Research Consortium. *Circulation*. 2011;123(23):2736–47.
 19. Fanaroff AC, Vora AN, Lopes RD. Non-vitamin K antagonist oral anticoagulants in patients with valvular heart disease. *Eur Hear Journal, Suppl*. 2022;24:A19–31.
 20. Yokoyama Y, Briasoulis A, Ueyama H, et al. Direct oral anticoagulants versus vitamin K antagonists in patients with atrial fibrillation and bioprosthetic valves: A meta-analysis. *J Thorac Cardiovasc Surg*. 2021;2401. Available from: <https://doi.org/10.1016/j.jtcvs.2021.07.034>
 21. Guimarães PO, Pokorney SD, Lopes RD, et al. Efficacy and safety of apixaban vs warfarin in patients with atrial fibrillation and prior bioprosthetic valve replacement or valve repair: Insights from the ARISTOTLE trial. *Clin Cardiol*. 2019;42(5):568–71.
 22. Patel M, Mahaffey K, Garg J, et al. Rivaroxaban versus Warfarin in Nonvalvular Atrial Fibrillation -

- The ROCKET-AF trial. *N Engl J Med*. 2011;365:687–96.
23. Hellenbart E, Faulkenberg K, Finks S. Evaluation of bleeding in patients receiving direct oral anticoagulants. *Vasc Health Risk Manag*. 2017;13:325–42. Available from: <http://dx.doi.org/10.2147/VHRM.S121661>
 24. Granger CB, Lopes RD, Hanna M, et al. Clinical events after transitioning from apixaban versus warfarin to warfarin at the end of the Apixaban for Reduction in Stroke and Other Thromboembolic Events in Atrial Fibrillation (ARISTOTLE) trial. *Am Heart J*. 2015;169(1):25–30. Available from: <http://dx.doi.org/10.1016/j.ahj.2014.09.006>
 25. Franchi F, Rollini F. Switching Oral Anticoagulant Therapy in Patients With Atrial Fibrillation Undergoing Percutaneous Coronary Intervention. *JACC Cardiovasc Interv*. 2019;12(23):2342–5.
 26. Granger CB, Alexander JH, McMurray JJ V, et al. Apixaban versus Warfarin in Patients with Atrial Fibrillation. *N Engl J Med*. 2011;365:687–96.
 27. Verstraete A, Herregods MC, Verbrugghe P, et al. Antithrombotic Treatment After Surgical and Transcatheter Heart Valve Repair and Replacement. *Front Cardiovasc Med*. 2021;8(August):1–12.
 28. Shenthari J. Management of atrial fibrillation in rheumatic heart disease. *Heart Rhythm O2*. 2022;3(6):752–9. Available from: <https://doi.org/10.1016/j.hroo.2022.09.020>

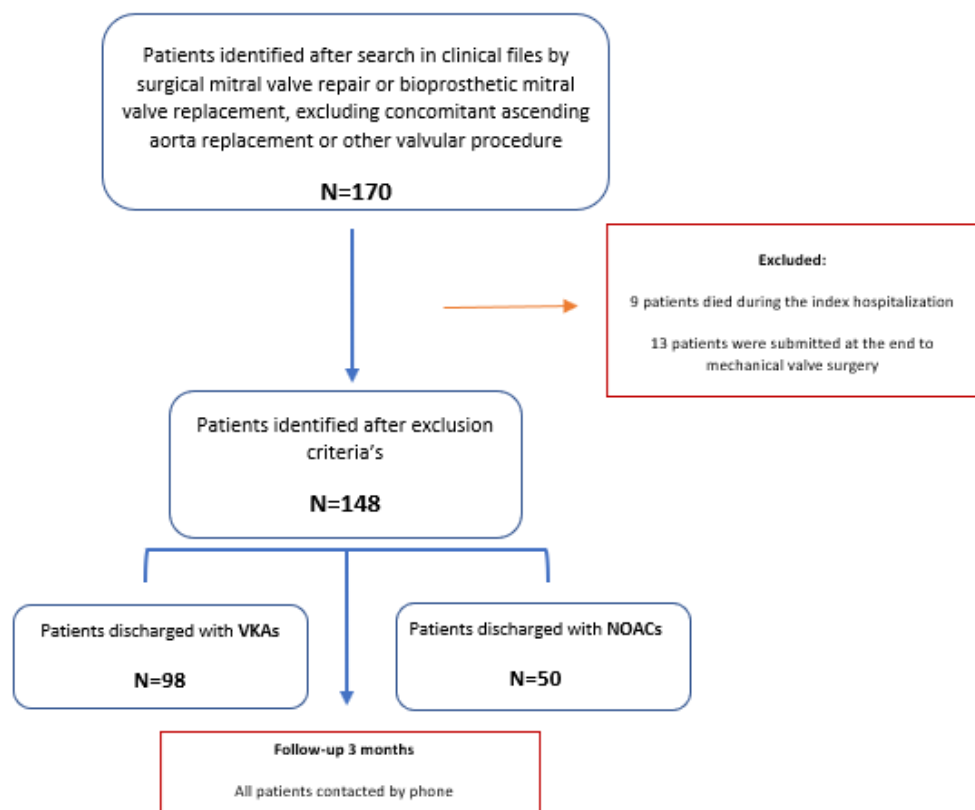


Figure 1 Study flowchart. NOACs: non-vitamin K antagonist oral anticoagulants; VKAs: vitamin K antagonists.

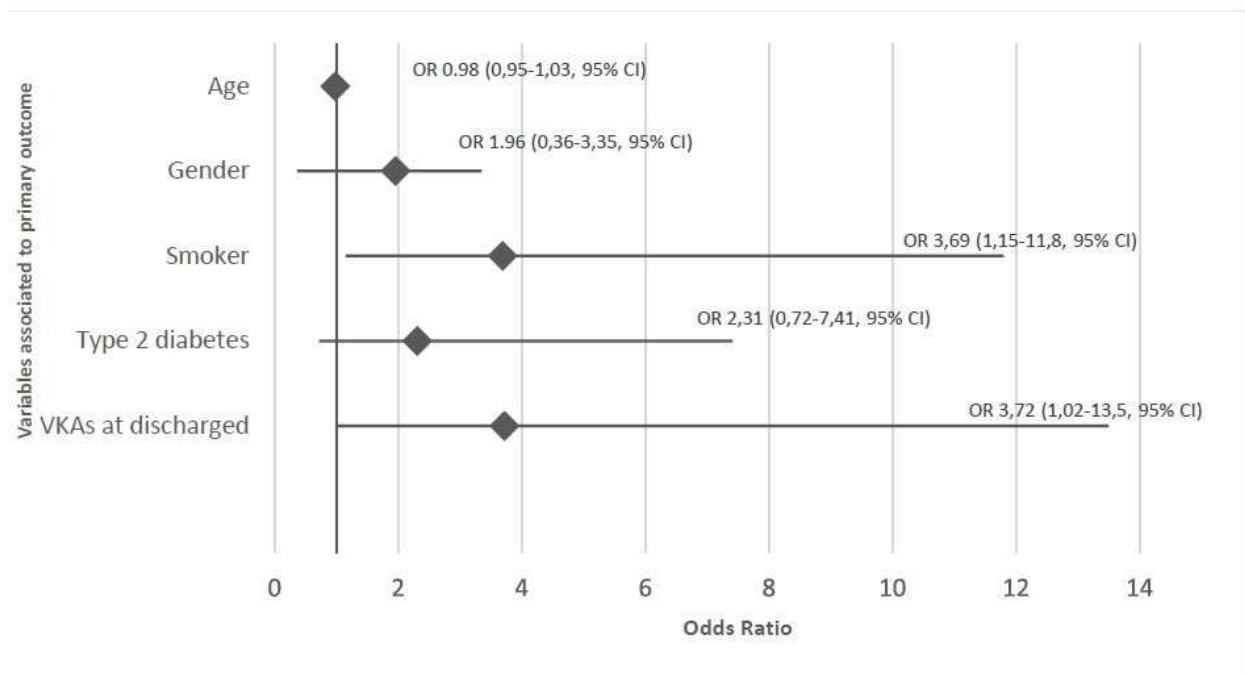


Figure 2 Multivariate analysis (logistic regression) for the primary outcome. CI: confidence interval; OR: odds ratio; VKAs: vitamin K antagonists.