

The Value of an Intervention for Improving Secondary Prevention in Patients Undergoing Cardiac Surgery

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Given that treatment for secondary prevention in patients undergoing cardiac surgery is underused, we devised a hospital intervention to increase its implementation. The intervention involved all physicians in the department of cardiac surgery agreeing to complete a report on each patient before hospital discharge. The document recorded the indications for the recommended treatments, and prompted for details of the drugs prescribed, the doses used, the reasons for not prescribing the recommended drugs, if that was the case, and the use of alternative medicines. The efficacy of the intervention was evaluated by comparing the rate of drug use in the year in which it was introduced (2003, n=341) with retrospective data on the rate in the previous year (n=369). The rates of use of aspirin, statins, angiotensin-converting enzyme inhibitors, and beta-blockers by patients who required them all showed an absolute increase, of 13.4%, 38.3%, 21.8%, and 21.5%, respectively. In conclusion, the introduction of a simple and inexpensive intervention was able to significantly increase the use of drugs for secondary prevention in patients undergoing cardiac surgery.

Key words: Secondary prevention. Cardiac surgery. Quality improvement.

Evaluación de una intervención para mejorar la prevención secundaria en pacientes sometidos a cirugía cardíaca

Dada la infrautilización de los tratamientos de prevención secundaria en pacientes sometidos a cirugía cardíaca, se diseñó una intervención hospitalaria para mejorar su empleo consistente en el compromiso de los miembros del servicio de cirugía cardíaca de cumplimentar antes del alta hospitalaria un formulario que recordaba las indicaciones de los tratamientos recomendados, preguntaba por su prescripción, la dosis empleada, la causa de no prescribir, si era el caso, y el uso de fármacos alternativos. Su eficacia se evaluó comparando la tasa de utilización de los fármacos el año de su uso, 2003 (n = 341), con la del año previo, obtenida retrospectivamente (n = 369). El uso de ácido acetilsalicílico, estatinas, inhibidores de la convertasa angiotensínica y bloqueadores beta en candidatos ideales aumentó en total el 13,4, el 38,3, el 21,8 y el 21,5% respectivamente. En conclusión, una intervención sencilla y barata fue capaz de mejorar significativamente el empleo de fármacos de prevención secundaria en pacientes sometidos a cirugía cardíaca.

Palabras clave: Prevención secundaria. Cirugía cardíaca. Mejora de calidad.

INTRODUCTION

Patients who are referred for cardiac surgery are increasingly older and present with a high prevalence of cardiovascular risk factors.^{1,2} Nevertheless, a number of nonsurgical measures for secondary prevention are reportedly underused, both in cardiovascular surgery departments^{3,4} and other settings.⁵⁻⁸ For this reason, we

decided to carry out a study to analyze the magnitude of the problem and evaluate the efficacy of an intervention focusing on improving prescription of secondary cardiovascular prevention measures in patients who undergo cardiac surgery.

METHODS

We reviewed the pharmacological measures for secondary cardiovascular prevention recommended in the major clinical practice guidelines.⁹⁻¹⁴ The conclusions compiled are summarized in Table 1.

An agreement was reached by which no patient would be discharged from the hospital in 2003 without the

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TABLE 1. Compilation of the Indications for Secondary Prevention Measures

Acetylsalicylic acid	Coronary artery surgery, coronary artery, cerebrovascular, or peripheral vascular disease Biological prosthesis in the absence of high thrombotic risk (3 months)
Statins	Cardiovascular disease plus one or more of the following: prior hypercholesterolemia or TC >200 mg/dL, LDL >100 mg/dL, TG >250 mg/dL DM2 plus another CVRF plus one or more of the following: LDL >100 mg/dL, TG >200 mg/dL DM without other CVRF plus one or more of the following: LDL >130 mg/dL, TG >250 mg/dL 2 CVRF or HDL <35 mg/dL plus one or more of the following: TC >200 mg/dL, LDL >100 mg/dL, TG >250 mg/dL No cardiovascular disease or DM plus one or more of the following: CT >240 mg/dL, LDL >130 mg/dL, TG >250 mg/dL
ACE inhibitors	LVEF <0.4 or LVEF <0.5 and heart failure History of myocardial infarction and changes in contractility DM
Beta-blockers	Ischemic heart disease Compensated heart failure or ventricular dysfunction of any cause
Oral anticoagulants	Mechanical valve prosthesis Biological valve prosthesis plus one or more of the following: atrial fibrillation, ventricular dysfunction, previous thromboembolism, hypercoagulability Atrial fibrillation and structural heart disease or history of embolism

ACE indicates angiotensin-converting enzyme; CVRF, cardiovascular risk factor; DM, diabetes mellitus; HDL, high-density lipoproteins; LVEF, left ventricular ejection fraction; LDL, low-density lipoproteins; TC, total cholesterol; TG, triglycerides.

completion of a form (Figure 1) that included personal details, diagnosis, type of surgery, cardiovascular risk factors, and personal history of each. In addition, any drug treatment prescribed for cardiovascular prevention should be recorded. The document made it necessary to review, on an individual basis, the indication and dosage of each drug or, should it be the case, the reason for which it had not been prescribed. A candidate was defined as that patient in whom a drug was indicated, according to the clinical practice guidelines, and an optimal candidate as a patient with indication and no contraindications. Subsequently, a study was designed to compare the group of patients who were discharged during 2003 with a control group consisting of the patients who were discharged in 2002, for whom the form was completed retrospectively.

Statistical Analysis

We carried out a descriptive analysis of the characteristics of the patients in each group to assess their homogeneity. The 2 groups were then compared in terms of prescription rates, adjusted to the different prevention measures employed in each. Student *t* test was used for the comparison of 2 means and ANOVA for the comparison of more than 2 means. The proportions were analyzed by the χ^2 test.

RESULTS

A total of 710 patients were recruited (369 in 2002 and 341 in 2003). Table 2 shows the clinical characteristics of each group. The rates of utilization of the secondary

prevention treatments in the 2 groups (2002/2003) are given below.

Figure 2 summarizes the percentage of optimal candidates in whom the corresponding secondary prevention measures were applied before and after the intervention.

Treatment with acetylsalicylic acid was received by 81.8% of the optimal candidates in 2002 and by 95.2% in 2003 ($P=.15$).

The number of patients who were optimal candidates for statin therapy was 203 (55%) in 2002 and 228 (66.8%) in 2003. Of these, 104 (51.2%) and 195 (97%), respectively, received treatment with statins ($P<.001$).

In 2002, 97.8% of the patients were optimal candidates for treatment with angiotensin-converting enzyme (ACE) inhibitors, whereas the percentage for 2003 was 92.3%; of these, 75.5% received ACE inhibitors in 2002 and 97.3% in 2003 ($P<.001$).

With respect to treatment with beta-blockers, 89% were optimal candidates in 2002 and 89.9% in 2003; of these patients, 73.6% received these drugs in 2002 and 95.2% in 2003 ($P<.001$).

All the optimal candidates for oral anticoagulation therapy in the 2 groups received this treatment.

DISCUSSION

The present study demonstrates that a simple and inexpensive intervention, such as the agreement of the members of a department to complete a reminder form prior to the discharge of each patient, can improve the use of secondary prevention drug treatments in patients who have undergone cardiac surgery.

Name _____		Sex _____		Age _____	
File no. _____		Date of birth ____/____/____		Date of admission ____/____/____	
				Date of discharge ____/____/____	
Diagnosis _____		Type of CVS _____			
CVRF	HT <input type="radio"/> Yes <input type="radio"/> No Smoking <input type="radio"/> Yes <input type="radio"/> No Hyperlipidemia <input type="radio"/> Yes <input type="radio"/> No Diabetes <input type="radio"/> Yes <input type="radio"/> No Fam hist CVD <input type="radio"/> Yes <input type="radio"/> No				
CV history	Coronary dis <input type="radio"/> Yes <input type="radio"/> No AMI <input type="radio"/> Yes <input type="radio"/> No Type AMI <input type="radio"/> Q-wave <input type="radio"/> Non-Q-wave Date 1st AMI ____/____/____				
	Cerebrovasc dis. <input type="radio"/> Yes <input type="radio"/> No PVD <input type="radio"/> Yes <input type="radio"/> No ICHF <input type="radio"/> Yes <input type="radio"/> No LVEF <50% <input type="radio"/> Yes <input type="radio"/> No AF <input type="radio"/> Yes <input type="radio"/> No				
Studiess	LVEF measured <input type="radio"/> Yes <input type="radio"/> No LVEF <input type="radio"/> >50 <input type="radio"/> 40-49 <input type="radio"/> 30-39 <input type="radio"/> <30				
	Lipids determined <input type="radio"/> Yes <input type="radio"/> No Cholesterol _____ mg/dL LDL _____ mg/dL HDL _____ mg/dL Triglycerides _____ mg/dL				
Nonpharmacological measurements Non-smoker <input type="radio"/> Yes <input type="radio"/> No Dietary <input type="radio"/> Yes <input type="radio"/> No Physical activity <input type="radio"/> Yes <input type="radio"/> No					
Pharmacological measurements					
ASA	<input type="radio"/> Yes → ASA dose _____ mg/day				
	<input type="radio"/> No → Reasons no ASA _____ Contraindications for ASA <input type="radio"/> Yes <input type="radio"/> No Alternative to ASA <input type="radio"/> Yes <input type="radio"/> No Clopidogrel <input type="radio"/> Yes <input type="radio"/> No				
Indications: Any patient with coronary artery surgery; coronary artery, cerebrovascular or peripheral arterial disease DM type 2 + another CVRF in the absence of high risk of bleeding Biological prosthesis in the absence of high risk (3 months)					
Statins	<input type="radio"/> Yes → Type of statin _____ Statin dose mg/dL _____ mg/dL				
	<input type="radio"/> No → Reasons no statins _____ Contraindication statins <input type="radio"/> Yes <input type="radio"/> No				
Indications: Cardiovascular disease + (previous hypercholesterolemia or TC >200 mg/dL or LDL >100 mg/dL or TG >400 mg/dL) DM type 2 + another CVRF + (LDL >100 mg/dL or TG >200 mg/dL or isolated DM + (LDL >130 mg/dL or TG >400 mg/dL) Patient with (2 CVRF or HDL <35 mg/dL) + (TC >200 mg/dL or LDL >100 mg/dL or TG >400 mg/dL) No CVD or DM with (TC >240 mg/dL or LDL >130 mg/dL or TG >400 mg/dL)					
ACE	<input type="radio"/> Yes → Type of ACE inhibitor _____ ACE inhibitor dose _____ mg/dL				
	<input type="radio"/> No → Reasons no ACE _____ Contraindications ACE inhibitors <input type="radio"/> Yes <input type="radio"/> No Alternative IACE inhibitors <input type="radio"/> Yes <input type="radio"/> No ARB-II <input type="radio"/> Yes <input type="radio"/> No Other VD _____				
Indications: All patients with LVEF <40% or LVEF <50% and CHF All patients with a history of AMI and changes in contractility All diabetic patients					
BB	<input type="radio"/> Yes → Type of BB _____ BB dose _____ mg/dL				
	<input type="radio"/> No → Reasons no BB _____ Contraindications BB <input type="radio"/> Yes <input type="radio"/> No Alternative BB _____				
Indications: All patients with ischemic heart disease Compensated heart failure or left ventricular dysfunction of any cause					
Sintrom	<input type="radio"/> Yes → IRecommended INR <input type="radio"/> 2-3 <input type="radio"/> 2.5-3.5 <input type="radio"/> 3-4				
	<input type="radio"/> No → Reasons no Sintrom _____ Contraindications for Sintrom <input type="radio"/> Yes <input type="radio"/> No Alternative to Sintrom _____				
Indications: Mechanical or biological prosthesis with CVRF (AF, left ventricular dysfunction, previous thromboembolism, or hypercoagulation) Patients with AF and structural heart disease or a history of embolism					

Figure 1. ACE indicates angiotensin-converting enzyme; AMI, acute myocardial infarction; ARB, angiotensin receptor blocker (angiotensin II receptor antagonist); ASA, acetylsalicylic acid; AF, atrial fibrillation; AMI, acute myocardial infarction; BB, beta-blocker; CHF, congestive heart failure; CV, cardiovascular; CVD, cardiovascular disease; CVRF, cardiovascular risk factor; CVS, cardiovascular surgery; DM, diabetes mellitus; HDL, high-density lipoprotein; HT, hypertension; INR, international normalized ratio; LDL, low-density lipoprotein; LVEF, left ventricular ejection fraction; PVD, peripheral vascular disease; TC, total cholesterol; TG, triglycerides; VD, vasodilator.

TABLE 2. Clinical Characteristics of the Patients Who Underwent Cardiac Surgery in 2002 and 2003

	2002 (n=369)	2003 (n=341)	P
Age, mean (SD), y	65.5 (11.7)	64.5 (12.3)	.31
Women, n (%)	140 (38.1)	134 (44.7)	.09
Hospital stay, mean (SD), d	19.5 (18.1)	21.7 (66)	.54
Hypertension, n (%)	191 (51.8)	153 (50.8)	.81
Smoking, n (%)	149 (40.1)	90 (29.9)	.005
Dyslipidemia, n (%)	162 (43.9)	142 (47.2)	.43
Diabetes mellitus, n (%)	97 (26.3)	67 (22.3)	.24
Family history, n (%)	43 (11.7)	38 (12.6)	.72
Arteriosclerosis, n (%)	123 (33.3)	87 (28.9)	.24
Previous myocardial infarction, n (%)	63 (17.1)	53 (17.6)	.92
Previous stroke, n (%)	39 (10.6)	13 (4.3)	.003
History of heart failure, n (%)	103 (27.9)	78 (25.9)	.6
Ejection fraction <0.4, n (%)	75 (20.3)	50 (16.6)	.23

SD indicates standard deviation.

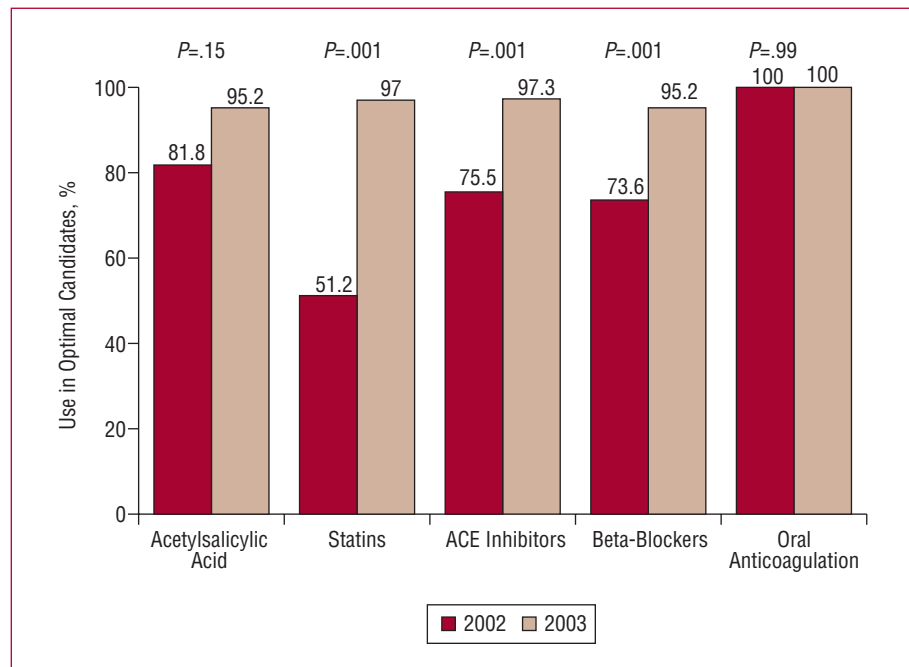


Figure 2. Percentage of optimal candidates who received the indicated treatment in 2002 and 2003. ACE indicates angiotensin-converting enzyme.

A number of secondary cardiovascular prevention measures have been shown to be capable of improving survival and reducing the incidence of new cardiovascular events.⁹⁻¹⁴ Despite the fact that cardiovascular prevention measures are employed effectively prior to surgery, the incidence of cardiovascular risk factors in patients referred for cardiac surgery remains high. A number of studies have been carried out to verify the degree of compliance with the recommendations of the clinical practice guidelines in cardiac patients. These studies have demonstrated both the underuse of these recommendations and the wide variability in their use, both in Spain^{5,6} and in other parts of the world.^{7,8} Although several reports have analyzed the processes related to the improved quality of patient management in the field of cardiology,¹⁵⁻²⁰

there is much less information on the patients who undergo cardiac surgery. We have managed to optimize the prescription of all the pharmacological measures, achieving a rate of compliance on the part of optimal candidates of nearly 95%. Our written form is an inexpensive, simple, highly manageable and reproducible tool that takes very little time to complete. It might be logical to think that, being subjected to a protocol to such a great extent, the tool should have resulted in a utilization rate of 100%; however, it must be taken into account that the introduction of a new tool in a department requires a certain amount of time and training.

One of the limitations of the study is that, since there is no nonintervention control group, the establishment of a cause-effect relationship may be questionable.

We consider that it would not have been ethical to fail to offer a system that was clearly going to improve the quality of care to all patients and, thus, we decided to use a historical control group.

In conclusion, a simple and inexpensive intervention, based on a manageable and reproducible document, is capable of producing a very substantial improvement in the use of secondary prevention methods in patients subjected to cardiac surgery.

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