

Management of Atrial Fibrillation after Cardiac Surgery

Essay

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Care Medicine*

By

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List of Abbreviation

<i>ACEIs</i>	Angiotensin –Converting Enzyme-Inhibitor
<i>ACS</i>	Acute Coronary Syndrome
<i>AF</i>	Atrial Fibrillation
<i>AFFRiM</i>	Atrial Fibrillation Follow up investigation of Rhythm Managment
<i>aPTT</i>	Activated Partial Thrombin Time
<i>ARB</i>	Angiotensin Receptor Blocker
<i>AV</i>	Atrio-Ventricular
<i>AVERRoES</i>	Reduce Apixban Versus acetylsalicylic acid to Reduce the Rate of Embolic Stroke in atrial fibrillation
<i>b.i.d</i>	bis in die(twice daily)
<i>BMS</i>	Bare Metal Stent
<i>CHA2DS2-VASC</i>	Cardiac failure, Hypertention, Age>75 (doubled), Diabetes, Stroke (doubled), Vascular disease, Age (65-74), Sex Category female
<i>CHADS2</i>	Cardiac failure, Hypertention, Age, Diabetes, Stroke (doubled)
<i>CrCl</i>	Creatinin Clearance
<i>CRT</i>	Cardiac Resynchronization Therapy
<i>CT</i>	Computerized Tomography
<i>CV</i>	Cardioversion
<i>DC</i>	Direct Cardioversion
<i>DCC</i>	Direct Current Cardioversion
<i>DIAMOND</i>	Danish Investigation of Arrhythmia and Mortality ON Dofetilide
<i>ECG</i>	Electrocardiograph
<i>ER</i>	Extended Release formulation
<i>GP</i>	Ganglionic Plexi
<i>HAS-BLED</i>	Hypertention, Abnormal renal or liver function, Stroke, Bleeding history, Labile INR, Elderly>65, Drug or alchol.

<i>I.V</i>	IntraVenous
<i>ICE</i>	Intra Cardiac Echocardiography
<i>ICH</i>	IntraCranial Hemorrhage
<i>INR</i>	International Normalized Ratio
<i>IVC</i>	Inferior Vena Cava
<i>LA</i>	Left Atrium
<i>LIPV</i>	Left Inferior Pulmonary Vein
<i>LMWH</i>	Low Molecular Weight Heparin
<i>LSPV</i>	Left Superior Pulmonary Vein
<i>LV</i>	Left Ventricle
<i>LVED</i>	Left Ventricular End Diastole
<i>LVEF</i>	Left Ventricular Ejection Fraction
<i>MI</i>	Myocardial Infarction
<i>MRI</i>	Magnetic Resonance Imaging
<i>N.A</i>	Not Applicable
<i>NA</i>	Neuroaxial Anesthesia
<i>NOACs</i>	Novel Oral Anticoagulants
<i>NYHA</i>	New York Heart Association
<i>o.d.</i>	Once daily
<i>OAC</i>	Oral Anticoagulant
<i>PAF</i>	Paroxysmal Atrial Fibrillation
<i>PCI</i>	Percutaneous Intervention
<i>PIAF</i>	Pharmacological Intervention in Atrial Fibrillation
<i>PPIs</i>	Proton Pump Inhibitors
<i>PT</i>	Prothrombin Time
<i>PT</i>	Prothrombin Time
<i>PV</i>	Pulmonary Vein
<i>PVPs</i>	Pulmonary Vein Potentials
<i>PVs</i>	Pulmonary Veins
<i>RACE</i>	Rate Control versus Electrical cardioversion for persistent atrial fibrillation
<i>RELY</i>	Randomized Evaluation of Long anticoagulation therapy with dabigatran
<i>RF</i>	RadioFrequency
<i>RIPV</i>	Right Inferior Pulmonary Vein

<i>ROCKET</i>	Rivoroxaban Once daily oral-direct factor Xa inhibition Compared with vit K antagonism for prevention of stroke and Embolism Trial in atrial fibrillation
<i>RRR</i>	Relative Risk Reduction
<i>RSPV</i>	Right Superior Pulmonary Vein
<i>STAF</i>	The Strategies of Treatment of Atrial Fibrillation
<i>SVC</i>	Superior Vena Cava
<i>TIA</i>	Transient Ischemic Attack
<i>TOE</i>	TransOesophageal Echo
<i>TTR</i>	Time in Theraputic Range
<i>UFH</i>	UnFractionated Heparin
<i>UH</i>	Unfractionated Heparin
<i>VKA</i>	Vitamin K Antagonist

Introduction

Cardiac dysrhythmias are common during anesthesia and surgery and occur in patients with structural heart disease or normal hearts. The aggravating factor is often physiologic imbalance unique to perioperative settings e.g. anesthetic or adjuvant drugs, adrenergic stress, acid-base and electrolyte imbalance (*Psaty et al., 2002*).

Absolute incidence rates for postoperative atrial fibrillation vary depending on many variables, including types of procedures, patient age and comorbidities, criteria for diagnosis and methods of ECG monitoring. When diagnosed based on intermittently obtained 12 lead ECGs. Atrial fibrillation is reported in 11% of patients compared with an incidence of > 40% when Holter monitoring is employed. With a reported incidence of up to 65% after open cardiac surgery (*Giovanni et al., 2010*).

Increased left atrial size due to valvular lesions, increased sympathetic nervous system activity or vagal tone, increased intravascular volume or perioperative hypovolemia, hypothermia, and potassium and magnesium deficiencies have all been associated with the development of atrial fibrillation (*Christians et al., 2001*).

Introduction

Coronary artery disease, cardiomegaly and premature atrial contractions were significantly associated with perioperative atrial arrhythmias in noncardiothoracic patients (*Jennifer et al., 2010*).

Postoperative atrial fibrillation usually tends to occur within 2 to 4 days after surgery, with a peak incidence on postoperative day 2. Although postoperative AF is usually a well-tolerated, transient problem, it is associated with the need for additional treatment, prolonged hospital stay, increased costs (*Halonen et al., 2010*).

Ventricular rate control is recommended in patients with AF without haemodynamic instability. Restoration of sinus rhythm by Direct Current Cardioversion, DCC, is recommended in patients who develop post-operative AF and are haemodynamically unstable. Current indications for curative ablation of AF are limited to patients who remain symptomatic despite the use of antiarrhythmic agents (*Vahanian et al., 2010*).

Aim of the Work

The aim of this work is to discuss the pathophysiology and up to date in management of atrial fibrillation postcardiac surgery.

Pathophysiology of Atrial Fibrillation

Definition:

Atrial fibrillation is a supraventricular tachyarrhythmia characterized by uncoordinated atrial activation with consequent deterioration of atrial mechanical function. On the electrocardio-gram (ECG), AF is characterized by the replacement of consistent P waves by rapid oscillations or fibrillatory waves that vary in amplitude, shape, and timing, associated with an irregular, frequently rapid ventricular response when atrioventricular conduction is intact (*Figure 1,2*) (*valentin fuster et al., 2011*).



Figure (1): Normal ECG (*valentin fuster et al., 2011*).



Figure (2): ECG showing atrial fibrillation (*valentin fuster et al., 2011*).

Pathophysiology of Atrial fibrillation

The ventricular response to AF depends on electrophysiological properties of the AV node and other conducting tissues, the level of vagal and sympathetic tone, the presence or absence of accessory conduction pathways, and the action of drugs. Regular cardiac cycles (R-R intervals) are possible in the presence of AV block or ventricular or AV junctional tachycardia (*Figure 3*) (*salvatore et al., 2008*).

In patients with implanted pacemakers, diagnosis of AF may require temporary inhibition of the pacemaker to expose atrial fibrillatory activity (*valentin fuster et al., 2011*).

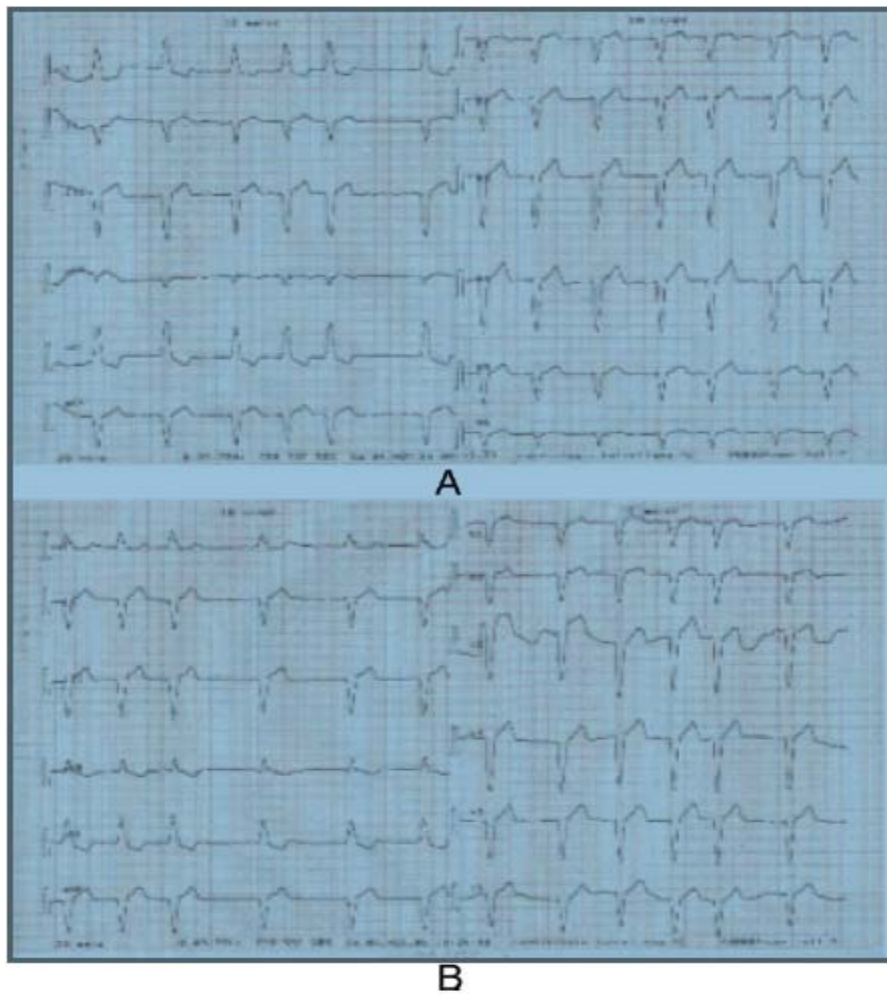


Figure (3): ECG showing atrial fibrillation with left bundle Branch block (*Salvatore et al.,2008*).

Classification:

Various classification systems have been proposed for AF. One is based on the ECG presentation (*Levy et al.,1998*). another is based on epicardial (*Allessie et al., 1994*), endocavitary recordings or noncontact mapping of atrial electrical activity. Several clinical classification schemes have also been proposed, but none fully accounts for all aspects of AF (*Levy, 2000*). To be clinically useful, a classification system must be based on a sufficient number of features and carry specific therapeutic implications. Assorted labels have been used to describe the pattern of AF, including acute, chronic, paroxysmal, intermittent, constant, persistent, and permanent (*Figure 4*). Although the pattern of the arrhythmia can change over time, it may be of clinical value to characterize the arrhythmia at a given moment (*Valentin fuster et al.,2001*).

The clinician should distinguish a first-detected episode of AF, whether or not it is symptomatic or self-limited, recognizing that there may be uncertainty about the duration of the episode and about previous undetected episodes.