have also suffered from a TTC, because there were no detailed data concerning cardiac function in these reports. In our report, two other trigger factors also could have promoted the TTC: the cesarean delivery itself or the oxytocin infusion, which induced a junctional tachycardia with myocardial ischemia. However, the delay between the stressful trigger and the first symptom is usually less than 2 h in the published cases of TTC after cesarean delivery.1 Moreover, our patient became transiently asymptomatic when the junctional tachycardia ended. These factors argue in favor of a probable relationship between sulprostone and the TTC.

The lesson gleaned from the described cases is worthwhile: cardiac symptoms, LV systolic dysfunction, or a slight troponin increase after sulprostone administration should lead to a suspicion of TTC. Moreover during cesarean delivery, the differential diagnosis could be an acute coronary syndrome caused by spontaneous coronary artery dissection. These two diseases can be distinguished by TTE and coronary angiography. Clinicians should be aware of this potential adverse effect when monitoring patients receiving sulprostone.

Pierre-Yves Courand,a,* Sophie Jenck,a Emmanuel Cassar,a Emmanuelle David,b Nicolas Carrabin,d and Pierre Lantelmea

aService de Cardiologie, Hôpital de la Croix-Rousse, Hospices Civils de Lyon, Lyon, France
bService d’Anesthésie-Réanimation, Hôpital de la Croix-Rousse, Hospices Civils de Lyon, Lyon, France
cService d’Obstétrique, Hôpital de la Croix-Rousse, Hospices Civils de Lyon, Lyon, France

*Corresponding author:
E-mail address: pycourand@hotmail.com (P.-Y. Courand).
Available online 24 February 2012

REFERENCES


Staphylococcus Aureus Endocarditis on Transcatheter Aortic Valves

Endocarditis por Staphylococcus aureus sobre válvula aórtica percutánea

To the Editor,

Transcatheter aortic valves are becoming an alternative for patients with symptomatic severe aortic stenosis when comorbidity makes conventional valve replacement surgery an unaffordable risk. We describe one of the first cases of infective endocarditis on the CoreValve® model (Medtronic, Minneapolis, Minnesota) prosthetic aortic valve.

An 81-year-old male with dyspnea secondary to severe degenerative aortic stenosis. Comorbidity consisted of diabetes mellitus, chronic kidney failure, and a severely depressed left ventricular ejection fraction with pulmonary hypertension >60 mmHg. The coronary arteries showed no significant lesions. The surgical risk according to EuroSCORE was 29%, so it was decided to implant a CoreValve® transcatheter prosthesis, which took place after prophylaxis with ampicillin, with no major complications. According to the aortography, there was moderate, grade II/IV residual aortic regurgitation and the patient was discharged after a week.

During the following months, an improvement of 50% was seen in the left ventricular systolic function, with a decrease of pulmonary pressure to 45 mmHg. The patient remained asymptomatic, except for an admission 4 months after the implant for self-limiting gastrointestinal bleeding leading to anemia. A colonoscopy was performed without prior antibiotic prophylaxis and diverticulosis was diagnosed.
Two months after the colonoscopy, the patient was admitted with 2-week symptoms of discomfort, disorientation, fatigue, fever, cough, and dyspnea. The physical examination revealed basal crackles and normal prostatic sounds on auscultation. There was no hepatomegaly, splenomegaly, or skin lesions. The laboratory data showed systemic infection. The electrocardiogram showed left bundle branch block with a prolonged PR interval, similar to immediately after implantation of the prosthesis. The chest X-ray showed a diffuse interstitial pattern without clear condensation.

With the diagnosis of sepsis of probable pulmonary origin, blood cultures were taken and treatment started with cefazidine and ciprofloxacin. The patient’s clinical condition worsened with persistent fever and a decreased level of consciousness. A growth of methicillin-sensitive Staphylococcus aureus was observed in all 3 blood cultures. Transthoracic (Fig. 1) and transesophageal (Fig. 2 and Video) echocardiograms were performed that, despite the interference with the metal prosthetic mesh, showed a vegetation of 0.8 cm maximum diameter and area of 0.3 cm² attached to the aortic prosthesis, with no significant failure or signs of periannular complications. A 6-week treatment of cloxacillin and rifampicin was started, including gentamicin in the initial 2 weeks, following the current guidelines for treatment of staphylococcal prosthetic endocarditis. A computed tomography scan ruled out stroke. Later evolution was satisfactory, with disappearance of fever and an improved level of consciousness. At discharge the patient was asymptomatic, with no systemic infection evidenced in laboratory tests, negative blood cultures, and an echocardiogram showing no vegetation, with slight transprosthetic aortic insufficiency.

This was one of the first published cases of transcatheter aortic valve endocarditis. Patients eligible for transcatheter aortic valve replacement are at a higher risk of infection from these devices, given their comorbidity. The femoral and transapical access may be a gateway for the microorganism. However, in our case it was more probably related to the later admission, given that almost 6 months had passed from the prosthesis implant to onset of symptoms, and to the expected aggressiveness of any prosthetic staphylococcal infection. However, it must be stressed that most microorganisms associated with endocarditis after colonoscopy are enteroccci or Streptococcus bovis. Given the symptoms, surgery was considered as a treatment and further endorsed by the improvement in left ventricular function and the disappearance of severe pulmonary hypertension, which reduced the operating risk. There are several cases of successful surgery within 30 days of a transcatheter valve implant, but only one within 6 months. Endothelialization of the prosthesis may have led to replacement of both the valve and aortic root, which would have increased the complexity of the surgery and surgical risk. This consideration, coupled with the favorable evolution of the medical treatment, led us to opt for conservative management.

Given the increasing use of transcatheter implantation, new cases of endocarditis on these devices will be seen. Maximizing asepsis and antibiotic prophylaxis during the implantation and in subsequent invasive procedures will be essential to minimizing the number of cases. The fragility of the recipients poses new diagnostic and treatment challenges. Records will need to be kept to monitor the peculiarities of this new phenomenon.

**FUNDING**

This case was partially funded by the Red de Centros Cardiovasculares, which is in turn funded by the Instituto de Salud Carlos III, Madrid, Spain.
SUPPLEMENTARY MATERIAL

Management of a Hypersensitivity Reaction to Thienopyridines: Prasugrel-Induced Fever and Hepatitis Resolved After Switching to Clopidogrel

Abordaje de una reacción de hiper sensibilidad a tienopiridinas: fiebre y hepatitis inducidas por prasugrel resueltas tras su sustitución por clopidogrel

To the Editor,

Approximately 1% of patients in treatment with clopidogrel experience type I or IV hypersensitivity reactions consisting, in general, of rash or hematologic disorders (thrombocytopenia or neutropenia). More exceptionally, there have been reports of liver toxicity and systemic inflammatory response syndrome. Recently, it has been proposed to replace this drug with prasugrel, a new third-generation thienopyridine, in situations in which antiplatelet therapy cannot be discontinued and it is therefore impossible to apply desensitization protocols. To our knowledge, there are no reports in the literature of the reverse procedure in which a serious hypersensitivity reaction to prasugrel resolves after discontinuation of the drug and initiation of treatment with clopidogrel.

A 40-year-old woman with a history of smoking was attended in a center other than our own for oppressive precordial chest pain. After a positive result in the ischemia detection test, a coronary angiogram was performed. This showed severe lesions in the left anterior descending artery and the diagonal coronary artery; revascularization was achieved by direct stenting, with a good initial angiographic outcome. At the time, antiplatelet therapy with acetylsalicylic acid was initiated (100 mg/day) along with clopidogrel (300 mg loading dose). A few hours after the procedure, the patient developed acute pulmonary edema with ST segment elevation in the anteroseptal leads. A repetition of the coronary angiogram showed complete acute thrombosis of the stent in the left anterior descending artery, and so the angioplasty procedure was repeated and clopidogrel was replaced by prasugrel (loading dose of 60 mg followed by 10 mg/day). Abciximab was added. Cardiogenic shock required orotracheal intubation and mechanical ventilation, as well as hemodynamic support with vasoactive drugs and intra-aortic balloon contrapulsation. The patient was then transferred to our hospital. On admission, the patient had a body temperature of 38.1 °C, a blood pressure of 120/60 mmHg, and a heart rate of 65 bpm. In the laboratory tests, of note were leukocytes 7.23 × 10^9/L (eosinophils 2.4%); aspartate transaminase 208 U/L; alanine transaminase 93 U/L; gamma glutamyl transpeptidase 240 U/L; and alkaline phosphatase 170 U/L. In addition to the vasoactive drugs and antiplatelet agents, treatment included ranitidine, sodium heparin, furosemide, and eplerenone. After initiating infusion of levosimendan, the hemodynamic status of the patient progressively stabilized, and she was extubated and the vasoactive drugs and intra-aortic balloon contrapulsation were withdrawn. At the same time, there was a transient improvement in her liver function tests. However, despite starting empirical antibiotic therapy with meropenem and linezolid, fever persisted in the days that followed, with small daily oscillations in temperature (<1 °C) and transient remission after administration of antipyretics. Microbiological study were sterile or negative; these included serial blood cultures, urine cultures, and cultures of endotracheal aspirate; serology for hepatotropic viruses and the human immunodeficiency virus cytomegalovirus antigenemia; and the Mantoux skin-prick test. Similarly, full autoimmune testing was negative. The fever did not remit after changing venous and arterial catheters or suspending various drugs (ranitidine, heparin, and eplerenone). After 12 days, the number of stools began to increase (6-8 per day), with no pathological contents and also with negative etiologic study (copiculture and detection of Clostridium difficile toxin). Both the leukocyte count and acute-phase reactants (C-reactive protein and procalcitonin) remained within normal limits at all times, although there was a progressive change in the liver function tests (day 23 after admission: aspartate transaminase 115 U/L, alanine transaminase 155 U/L, and gamma glutamyl transpeptidase 723 U/L), and relative eosinophilia appeared (day 19, eosinophils 9.7%). Abdominal ultrasound did not reveal any liver or bile duct abnormalities. Finally, faced with suspected drug-mediated hypersensitivity reaction, it was decided to suspend prasugrel and start clopidogrel without making any other changes to therapy. In the first 24 h after the change, the fever and diarrhea remitted, and there was a progressive return to normal liver function tests and leukocyte differentials (Figure). After 7 days, the patient had a second episode of low cardiac output that required administration of vasoactive drugs. This time, there were no accompanying abnormal laboratory results or fever. As she had a left ventricular ejection fraction of 5% to 10%, with no possibility of revascularization, she was included on the waiting list for heart transplantation. This procedure was performed on day 32 after admission, without the reappearance of symptoms in the

REFERENCES


doi:10.1016/j.rec.2011.11.004