distribution in the methods section. We acknowledge that the inference that “a DHCA + RCP period of up to 80 minutes under a nasopharyngeal temperature of 18°C is safe” in our experience is rather anecdotal and cannot be totally substantiated. However, if we divide the patients into 2 groups—those who had DHCA + RCP for less than 60 minutes and those in whom this period lasted 60 minutes or more—we cannot find any difference in the incidence of mortality, stroke, and transient neurologic dysfunction. However, it is obvious that we should reduce the period of DHCA as much as possible, even in patients who require a complex repair of the aortic arch.

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Reconstruction of the aortic valve with autologous pericardium: An experimental study

To the Editor:

Valve repair has several advantages over prosthetic valve replacement, including low morbidity and mortality and lower risk of thromboembolism, hemorrhage, and septic endocarditis. Different techniques of mitral valve repair have been described, with good results. Unfortunately, the results of aortic valve repair are not as favorable. In this report I describe a new technique of aortic valve reconstruction with an autologous pericardial patch, which my colleagues and I have used.

In 5 cadaver hearts, autologous pericardium was fixed in 0.6% glutaraldehyde solution for 10 minutes. An aortotomy was used. Pericardium was cut to simulate the dimensions of the valve to be repaired on the basis of the concept of the aortic root geometry as a truncated cone. The diameter at the highest point of attachment of the leaflets (sinotubular diameter) is about 20% less than the diameter at the inlet (surgical anulus diameter). During systole the sinotubular diameter increases while the inlet diameter decreases, changing the root geometry from conical to cylindrical. The reverse occurs in diastole, at which time the leaflets tilt toward the ventricle.

The noncoronary cusp is fashioned first. It is measured at its attachment line and height with a soft wire (Fig 1, A) and the measurements are marked on the pericardium (Fig 1, B). The resulting “a-b” is the length at the sinotubular circumference but not the true length of the cusp. Therefore, we lengthened it to the size of its projection into the surgical (inlet) circumference.

Fig 1. Cusp size measurements and fashioning of the pericardial patch. A, Taking of the noncoronary cusp attachment line and height measurements with the soft wire. B, Marking of the sizes on the pericardium and fashioning of the pericardial patch.
The prerequisite to our method is the absence of significant distortion of aortic root geometry in hearts with severely diseased valves.

We wish to pursue this study further and look forward to collaborating with others in this endeavor.

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**REFERENCES**


**A stable model of chronic bilateral ventricular insufficiency (dilated cardiomyopathy) induced by arteriovenous anastomosis and doxorubicin administration in sheep**

To the Editor

Congratulations to Toyoda, Okada, and Kashem\(^1\) on their recent article, “A Canine Model of Dilated Cardiomyopathy Induced by Repetitive Intracoronary Doxorubicin Administration.” We need such a simple and reproducible animal model of stable, nonreversible chronic heart failure if we are to evaluate new surgical treatments for pre–end-stage congestive heart failure. The authors described cannulation of the coronary artery via the femoral artery and transcatheter infusion of doxorubicin, which resulted in visible signs of left ventricular insufficiency after 5 weekly doses of intracoronary doxorubicin; however, they reported that “the right ventricle seemed to be affected only slightly by doxorubicin.” Especially for cardiomyopathy research, one needs to create a model of bilateral heart failure.

In a recently completed (unpublished) investigation involving adult sheep, we created a bilateral model through first inducing right ventricular insufficiency by a surgical anastomosis between the right jugular vein and right external cardiac artery. After 1 month, we observed initial signs of insufficiency in the right ventricle (Table I) but not in the left. At this point, we began daily administration of (intravenous, not intracoronary) doxorubicin, which continued for the next 4 weeks and resulted in left ventricular insufficiency (Table II).

Please note that we made no attempt to determine whether the anastomosis or doxorubicin had the greater effect in inducing right ventricular insufficiency, although Toyoda, Okada, and Kashem observed that 5 weekly doses of the drug were insufficient for the purpose. On the other hand, Smink and colleagues\(^2\) induced quite stable bilateral ventricular insufficiency after an anastomosis was in place for only 3 months.