



Medline <1966 to July Week 4 2000>

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Citation 1.

Unique Identifier

98414735

Authors

Wernly JA. Crawford MH.

Institution

Division of Thoracic and Cardiovascular Surgery, University of New Mexico School of Medicine, Albuquerque, USA.

Title

Choosing a prosthetic heart valve. [Review] [59 refs]

Source

Cardiology Clinics. 16(3):491-504, 1998 Aug.

Abstract

Although most of the available prosthetic heart valves function remarkably well, the variety of available choices attests to the inability of any single one to fulfill the requirements of the ideal valve substitute. The mechanical prostheses include the caged-ball, tilting-disc, and bileaflet valves. Tissue valves available in the United States are the Carpentier-Edwards and Hancock porcine heterograft valves and the Carpentier-Edwards pericardial valve. Review of several large comparative studies on valve performance reveals that the overall results with tissue and mechanical valves are about equal at the end of 10 years. The characteristics of each type of valve substitute dictate the selection of one prosthesis in preference to others for a particular patient. Mechanical prostheses are recommended for patients without contraindications for anticoagulants. Tissue valves are reserved for patients over 65 years of age or for patients in whom anticoagulation is contraindicated. Multiple other patient-related factors need to be considered in selecting the appropriate valve, including the psychosocial situation and patient preference. [References: 59]

Citation 2.

Unique Identifier

98339145

Authors

Hirai S. Fukunaga S. Sueshiro M. Watari M. Sueda T. Matsuura Y.

Institution

First Department of Surgery, Hiroshima University School of Medicine, Japan.

Title

Assessment of a new silicone tri-leaflet valve seamlessly assembled with blood chamber for a low-cost ventricular assist device.

Source

Hiroshima Journal of Medical Sciences. 47(2):47-55, 1998 Jun.

Abstract

We have developed a practical, low-cost ventricular assist device (VAD) comprising a newly designed blood chamber with a silicone lenticular sac and two silicone tri-leaflet valves (STV), made en bloc. This new VAD is seamless, can be made cost-effectively and assembled with the blood chamber and valve as one body. This novel design should reduce the incident of thrombus formation because of the absence of a junction at the connecting ring and because of the use of flexible silicone materials which are biocompatible. In in vitro hemodynamics testing, a batch of 3 consecu

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Medtronic, Inc., Medtronic Vascular, Inc.,
& Medtronic Corevalve, LLC
v. Troy R. Norred, M.D.
Case IPR2014-00110



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