

This 510(k) summary of safety and effectiveness information is being submitted in accordance with the requirements of SMDA 1990 and 21 CFR 807.92.

The assigned 510(k) number is: K972570

**Applicant Information:**

Date Prepared: July 9, 1997  
Name: Heartport, Inc.  
Address: 200 Chesapeake Drive  
Redwood City, CA 94063

OCT - 1 1997

Contact Person: Marianne C. Drennan  
Phone Number: (415) 482-4405  
Fax Number: (415) 482-4346

**Device Information:**

Classification Class II  
Trade Name: Heartport® Introducer Sheath  
Common Name: Cardiovascular Surgical Devices - Catheter introducer  
21 CFR 870.1340

**Equivalent Devices:**

The subject device is substantially equivalent in intended use and method of operation to the currently marketed Heartport® Endoarterial Return™ Cannula.

**Intended Use:**

The Introducer Sheath with hemostasis valve is indicated for patients requiring introduction of catheters. It is intended for the hemostatic introduction and removal of vascular catheters such as the Heartport® Endoaortic Clamp™ Catheter.

**Non-Clinical Test Results:**

Performance testing of the subject device was not deemed necessary based on equivalence to the predicate device. The materials used in the subject device have proven biocompatibility.

**Summary:**

Based on the intended use, product performance and biocompatibility information provided in this notification, the subject device has been shown to be substantially equivalent to currently marketed predicate device.



Food and Drug Administration  
9200 Corporate Boulevard  
Rockville MD 20850

OCT - 1 1997

Ms. Marianne C. Drennan  
Heartport, Inc.  
200 Chesapeake Drive  
Redwood City, California 94063

Re: K972570  
Heartport® Introducer Sheath  
Regulatory Class: II (two)  
Product Code: 74 DYB  
Dated: August 20, 1997  
Received: August 22, 1997

Dear Ms. Drennan:

We have reviewed your Section 510(k) notification of intent to market the device referenced above and we have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

If your device is classified (see above) into either class II (Special Controls) or class III (Premarket Approval), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to-895. A substantially equivalent determination assumes compliance with the Current Good Manufacturing Practice requirement, as set forth in the Quality System Regulation (QS) for Medical Devices: General regulation (21 CFR Part 820) and that, through periodic QS inspections, the Food and Drug Administration (FDA) will verify such assumptions. Failure to comply with the GMP regulation may result in regulatory action. In addition, FDA may publish further announcements concerning your device in the Federal Register. Please note: this response to your premarket notification submission does not affect any obligation you might have under sections 531 through 542 of the Act for devices under the Electronic Product Radiation Control provisions, or other Federal laws or regulations.

Page 2 - Ms. Marianne C. Drennan

This letter will allow you to begin marketing your device as described in your 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801 and additionally 809.10 for in vitro diagnostic devices), please contact the Office of Compliance at (301) 594-4639. Additionally, for questions on the promotion and advertising of your device, please contact the Office of Compliance at (301) 594-4639. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21 CFR 807.97).

Other general information on your responsibilities under the Act may be obtained from the Division of Small Manufacturers Assistance at its toll-free number (800) 638-2041 or (301) 443-6597 or at its internet address "<http://www.fda.gov/cdrh/dsmamain.html>".

Sincerely yours,



Thomas J. Callahan, Ph.D.  
Director  
Division of Cardiovascular,  
Respiratory, and Neurological Devices  
Office of Device Evaluation  
Center for Devices and  
Radiological Health

Enclosure

**Indications for Use**

510(k) Number (if known): K972570

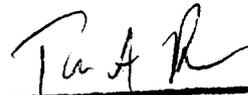
Device Name: Heartport® Introducer Sheath with Hemostasis Valve

**Indications for Use:**

The Introducer Sheath with hemostasis valve is indicated for patients requiring introduction of catheters. It is intended for the hemostatic introduction and removal of vascular catheters such as the Heartport® Endoaortic Clamp™ Catheter.

(PLEASE DO NOT WRITE BELOW THIS LINE - CONTINUE ON ANOTHER PAGE IF NEEDED)

Concurrence of CDRH, Office of Device Evaluation (ODE)



**(Division Sign-Off)**  
**Division of Cardiovascular, Respiratory,**  
**and Neurological Devices**  
510(k) Number K972570

Prescription Use              
(Per 21 CFR 801.109)

OR

Over- The Counter Use             
(Optional Format 1-2-96)