

K971019

## SUMMARY OF SAFETY AND EFFECTIVENESS

### CHASE VESSEL CANNULA

JUL 21 1997

#### I. General Information

- A. Generic Name: Vessel Cannula
- B. Trade Name of Device: Chase Vessel Cannula
- C. Applicant's Name and Address: CHASE MEDICAL, INC. , Richardson, TX
- D. Pre-market Notification Number: Not assigned

#### II. Indication for Use:

The Vessel Cannula is indicated for use during open-heart surgery to perfuse fluid into vein grafts.

#### III. Device Description:

The Vessel Cannula has stepped barbs at its tip and female luer fitting.

#### IV. Device Classification: Class I device

#### V. Safety and Effectiveness:

Substantial Equivalence: This device is substantially equivalent to the Cobe Labs Vessel Cannula (K760624).

#### VI. Other Safety and Effectiveness Data:

- Materials: All material are identical to the predicate device.
- Sterilization: Validated 100% Ethylene Oxide sterilization cycle (Overkill Method) SAL  $10^{-6}$

#### Functional Testing

All functional characteristics of the Chase Vessel Cannula are non-differentiable as compared with the predicate because both devices have the exact same fit, form, and material composition.

## SUMMARY OF SAFETY AND EFFECTIVENESS

Package Integrity:	Tyvek/Polymylar passed burst test per ASTM F1140-88
Shipping & Distribution Testing:	Per National Safe Transit Ass. vibration and drop tests
Accelerated Aging:	Two year shelf life



DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration  
9200 Corporate Boulevard  
Rockville MD 20856

Mr. Bert Davis  
President  
Chase Medical, Inc.  
1876 Firman Drive  
Richardson, Texas 75081

JUL 21 1997

Re: K971019  
Chase Vessel Cannula  
Regulatory Class: II (Two)  
Product Code: 74 DWF  
Dated: June 30, 1997  
Received: June 30, 1997

Dear Mr. Davis:

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.

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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) 594-4648. 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

510(k) Number (if known): K 971019

Device Name: \_\_\_\_\_

Indications For Use:

CHASE MEDICAL, INC.

VESSEL CANNULA

Intended Use:

The Vessel Cannula is indicated for use during open-heart surgery to perfuse fluid into vein grafts.

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

Concurrence of CDRH, Office of Device Evaluation (ODE)

*Ben R. Campbell*  
(Division Sign-Off)

Division of Cardiovascular, Respiratory,  
and Neurological Devices

510(k) Number K971019

Prescription Use X  
(Per 21 CFR 801.109)

OR

Over-The-Counter Use \_\_\_\_\_

(Optional Format 1-2-96)