K003901

JAN 1 7 2001

Confidential

Special 510(k) Device Modification
ANGIODYNAMICS Soft-Vu Sizing Catheter

General Information:

ANGIODYNAMICS, Inc. intends to introduce the following device into commercial distribution:

a)	**Trade Name:**	ANGIODYNAMICS Soft-Vu Sizing Angiographic Catheter
b)	**Legally Marketed Device:**	ANGIODYNAMICS Soft-Vu Hydrophilic Coated Angiographic Catheter
	510(k) Number:	K001578
c)	**Classification Name:**	Angiographic Catheter
d)	**Established Registration Number:**	1319211
e)	**Manufacturing Site Address:**	ANGIODYNAMICS, Inc. 603 Queensbury Avenue Queensbury, New York 12804
f)	**Sterilization Site Address:**	Griffith MicroScience Inc. 27 Park Rd. Glens Falls, NY 12801
g)	**Sterilizer Establishment Registration Number**	1319639
h)	**Classification:**	Regulation Number 870.1200 Product Code: DQO Regulatory Class II Cardiovascular Devices

i) **Device Equivalence:**

This product is substantially equivalent to the following device:

- Cook Royal Flush II Angiographic Sizing Catheter

This device does not present additional risks to patients or different considerations regarding safety and effectiveness than those presented by the predicate device.

j) **Performance Standards:** None Established

k) **ANGIODYNAMICS® Contact Information:**

Name:	Teri Juckett, Regulatory Affairs Associate
Address:	603 Queensbury Avenue Queensbury, New York 12804
Phone:	(518) 798-1215 extension 142
Fax:	(518) 798-3625

DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

JAN 17 2001

Ms. Teri Juckett
Regulatory Affairs Associate
AngioDynamics, Inc.
603 Queensbury Avenue
Queensbury, NY 12804

Re: K003901
Trade Name: AngioDynamics Soft-Vu Sizing Angiographic Catheter
Regulatory Class: II (two)
Product Code: DQO
Dated: December 14, 2000
Received: December 18, 2000

Dear Ms. Juckett:

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 legally marketed predicate 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 requirements, 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.

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) 594-4586. 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" (21CFR 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/dsma/dsmamain.html".

Sincerely yours,

For James E. Dillard III
Director
Division of Cardiovascular and
Respiratory Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

INDICATIONS FOR USE

510(k) Application: Special 510(k) Device Modification

Device Name: ANGIODYNAMICS Soft-Vu Sizing Catheter

Indications for Use:

The ANGIODYNAMICS Soft-Vu Sizing Catheter is for use where angiographic diagnosis is indicated.

Please do not write below this line - continue on another page if needed

Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use ✓ **OR** **Over-the-Counter Use** _____
(Per 21 CFR 801.109)

1/17/1

Division of Cardiovascular & Respiratory Devices
510(k) Number K003901