

K974769



**Allegiance Healthcare Corporation**  
1500 Waukegan Road  
McGaw Park, IL 60085  
847.473.1500  
FAX: 847.785.2461

JUN 12 1998

## SUMMARY OF SAFETY AND EFFECTIVENESS Appendix F

Manufacturer:	Allegiance Healthcare Corporation V. Mueller Business Unit 1435 Lake Cook Road Deerfield, IL 60015
Regulatory Affairs Contact	Patricia Sharpe-Gregg 1500 Waukegan Road McGaw Park, Illinois 60085
Telephone:	(847) 578-3636
Date Summary Prepared:	December 19, 1997
Product Trade Name:	Cosgrove™ Clamp
Common Name:	Vascular Clamp
Classification:	Vascular Clamp
Predicate Device: (K951413)	Allegiance Fogarty-Hydragrip® Surgical Clamp
Description:	The Allegiance Cosgrove™ Clamp is a reusable device which features a stainless steel handle, flexible shaft and jaws. The Cosgrove™ Clamp is designed to be used with a variety of inserts which gently surround blood vessels and offer differing degrees of atraumatic occlusion. Following the clamping of the blood vessel, the shaft of the device can be bent out of the way to enhance visualization of and access to the operative field.

Intended Use:

The Allegiance Cosgrove™ Clamp is intended to be used for the temporary occlusion of blood vessels and other delicate vessels during surgery. Surgical applications include pulmonary and gastrointestinal procedures, peripheral clamping, minimally invasive and standard open cardiovascular and cardio-thoracic procedures such as occlusion of the aorta and vena cava, cross-clamping of the aorta, etc.

Substantial Equivalence:

The Allegiance Cosgrove™ Clamp is substantially equivalent to the Fogarty-Hydragrip® Surgical Clamps and the Applied Vascular Atraumax™ Surgical Clamps in that:

- Intended use is the same
- Performance attributes are the same
- Atraumatic Jaw design characteristic is the same

Summary of Testing:

All materials used in the composition of the Cosgrove™ Clamp were subjected to performance and physical tests to evaluate the safety, effectiveness and reliability of the device. All test results were acceptable.



JUN 12 1998

Food and Drug Administration  
9200 Corporate Boulevard  
Rockville MD 20850

Ms. Patricia Sharpe-Gregg  
Manager, Regulatory Affairs  
Allegiance Healthcare Corporation  
1500 Waukegan Road, Bldg. MPWM  
McGaw Park, IL 60085

Re: K974769  
Trade Name: Cosgrove Clamp  
Regulatory Class: II  
Product Code: DXC  
Dated: May 30, 1998  
Received: June 14, 1998

Dear Ms. Sharpe-Gregg:

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 (Pre-market 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 pre-market 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) 594-4618. 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,



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

Enclosure



**Allegiance Healthcare Corporation**  
 1500 Waukegan Road  
 McGaw Park, Illinois 60085 USA  
 847-473-1500  
 FAX: 847-785-2461

**510(k) Notification Cosgrove™ Clamp**  
**V. Mueller Business Unit**  
 Page 1 of 1

**510(k) Number (if known):** Unknown

**Device Name:** Allegiance Cosgrove™ Clamp

**Indications For Use:** Used to occlude a blood vessel temporarily. Used in pulmonary and gastrointestinal procedures and can be used to clamp over indwelling catheters. Also used in minimally invasive and standard open cardiovascular procedures for temporary occlusion of a blood vessel.

(Division Sign-Off)  
 Division of Cardiovascular, Respiratory  
 and Neurological Devices  
 510(k) Number K974769

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

\_\_\_\_\_  
 Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use   
 (Per 21 CFR 801.109)

or Over-The Counter Use