



K103064

JAN 10 2011

510(k) SUMMARY

A summary of 510(k) safety and effectiveness information in accordance with 21 CFR 807.92.

SUBMITTER INFORMATION	
Name	CareFusion
Address	1500 Waukegan Road MPWM, McGaw Park, IL 60085 USA
Phone number	(847) 473-7404
Fax number	(847) 473-7990
Establishment Registration Number	1625685
Name of contact person	Joy Greidanus
Date prepared	10/28/2010
NAME OF DEVICE	
Trade or proprietary name	AVAmox Vertebral Balloon
Common or usual name	Inflatable Bone Tamp
Classification name	Primary: Arthroscope Secondary: Cement, Bone Vertebroplasty
Classification panel	Orthopedic
Regulation	Class II per 21CFR §888.1100, Procode HRX: Class II per 21CFR §888.3027, Procode NDN:
Product Code(s)	VBT1115, VBT1120
Legally marketed device(s) to which equivalence is claimed	CareFusion Inflatable Bone Tamp, K093463 Kyphx Inflatable Bone Tamp, K041454, K981251 Radiopaque Bone Cement, K043518
Reason for 510(k) submission	New Device
Device description	The Inflatable Bone Tamp (IBT) was designed for use in Balloon kyphoplasty. The balloon serves to create a cavity in the vertebral body, thereby reducing the fracture and preventing cement leakage, while still allowing for cement interdigitation. The balloon catheter is the functional part of the device that creates a cavity and reduces the fracture. The balloon catheter provides a conduit through which the physician can inflate the balloon at the distal end of the catheter. The wire stiffener provides stiffness to the balloon catheter to facilitate insertion through the access cannula
Intended use of the device	Intended for the reduction and fixation of fractures and/or creation of a void in cancellous bone in the spine for kyphoplasty (for use with CareFusion Radiopaque Bone Cement).



SUMMARY OF THE TECHNOLOGICAL CHARACTERISTICS OF THE DEVICE COMPARED TO THE PREDICATE DEVICE		
Characteristic	New Device	Predicate Kyphon IBT (K041454) CareFusion Vertebral Balloon (K093643)
Compatible cannula size	11G	10G
Balloon inflation medium	60% contrast recommended	60% contrast recommended
Balloon and catheter materials	Polyurethane	Polyurethane
Wire mandrel material	Stainless steel	Stainless steel
Balloon shape	Cylindrical	Kyphon: Variable CareFusion: Cylindrical
Maximum recommended inflation pressure	400 psi (27 ATM)	400 psi (27 ATM)
Maximum recommended inflation volume (15mm)	4 mL	Kyphon: 5 mL CareFusion: 4 mL
Maximum recommended inflation volume (20mm)	6 mL	Kyphon: No 20mm size Carefusion: 6 mL
PERFORMANCE DATA		
SUMMARY OF NON-CLINICAL TESTS CONDUCTED FOR DETERMINATION OF SUBSTANTIAL EQUIVALENCE		
Performance Test Summary-New Device		
Characteristic	Standard/Test/FDA Guidance	Results Summary
Inflation pressure	Constrained burst test	The 15mm and 20mm balloon catheters exceeded the requirements for the minimum burst pressure in a constrained environment
Inflation volume	Unconstrained burst test	The 15mm and 20mm balloon catheters exceeded the requirements for the minimum burst volume in an unconstrained environment
Balloon double wall thickness	Calibrated measurement	The double wall thickness of the 15mm and 20mm balloons was substantially equivalent to that of the predicate devices
SUMMARY OF CLINICAL TESTS CONDUCTED FOR DETERMINATION OF SUBSTANTIAL EQUIVALENCE AND/OR OF CLINICAL INFORMATION		
N/A – No clinical tests were conducted for this submission		
CONCLUSIONS DRAWN FROM NON-CLINICAL AND CLINICAL DATA		
The results of the non-clinical tests show that the CareFusion 11G balloon catheters meet or exceed all performance requirements, and are substantially equivalent to the predicate devices.		



Food and Drug Administration
10903 New Hampshire Avenue
Document Control Room -WO66-G609
Silver Spring, MD 20993-0002

CareFusion
% Ms. Joy Greidanus
Manager, Regulatory Affairs
1500 Waukegan Road
McGaw Park, Illinois 60085

JAN 10 2011

Re: K103064
Trade Name: Inflatable Bone Tamp
Regulation Number: 21 CFR 888.3027
Regulation Name: Polymethylmethacrylate (PMMA) bone cement
Regulatory Class: Class II
Product Code: NDN, HRX
Dated: December 13, 2010
Received: December 14, 2010

Dear Ms. Greidanus:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and 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) that do not require approval of a premarket approval application (PMA). 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. Please note: CDRH does not evaluate information related to contract liability warranties. We remind you, however, that device labeling must be truthful and not misleading.

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

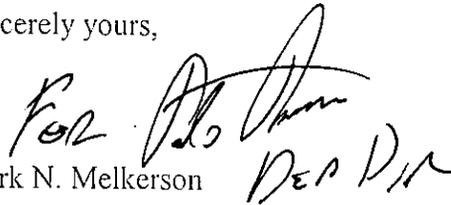
Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); medical device reporting (reporting of medical device-related adverse events) (21 CFR 803); good manufacturing practice requirements as set

forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801), please go to <http://www.fda.gov/AboutFDA/CentersOffices/CDRH/CDRHOffices/ucm115809.htm> for the Center for Devices and Radiological Health's (CDRH's) Office of Compliance. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21 CFR Part 807.97). For questions regarding the reporting of adverse events under the MDR regulation (21 CFR Part 803), please go to <http://www.fda.gov/MedicalDevices/Safety/ReportaProblem/default.htm> for the CDRH's Office of Surveillance and Biometrics/Division of Postmarket Surveillance.

You may obtain other general information on your responsibilities under the Act from the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (301) 796-7100 or at its Internet address <http://www.fda.gov/MedicalDevices/ResourcesforYou/Industry/default.htm>.

Sincerely yours,

A handwritten signature in black ink, appearing to read "FOR [Name] Dir", is written over the typed name and title.

Mark N. Melkerson
Director
Division of Surgical, Orthopedic,
and Restorative Devices
Office of Device Evaluation
Center for Devices and
Radiological Health



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Indication for Use

510(k) Number (if known): unknown at this time
Device Name: Inflatable Bone Tamp
Indications For Use: Intended for the reduction and fixation of fractures and/or creation of a void in cancellous bone in the spine for kyphoplasty (for use with CareFusion Radiopaque Bone Cement).

Prescription Use X or Over-The Counter Use _____

(Per 21 CFR 801.109)

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

Concurrence of CDRH, Office of Device Evaluation (ODE)



(Division Sign-Off)
Division of Surgical, Orthopedic,
and Restorative Devices

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