

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Activity Date 3-16-06
Publication Date 3-17-06
Author L. CLAWSON

[Docket Nos. 2005M-0435, 2005M-0475, 2005M-0473, 2005M-0478, 2005M-0454, 2005M-0399, 2005M-0477, 2005M-0476, 2005M-0492, 2005M-0474, 2005M-0504]

Medical Devices; Availability of Safety and Effectiveness Summaries for
Premarket Approval Applications

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is publishing a list of premarket approval applications (PMAs) that have been approved. This list is intended to inform the public of the availability of safety and effectiveness summaries of approved PMAs through the Internet and the agency's Division of Dockets Management.

ADDRESSES: Submit written requests for copies of summaries of safety and effectiveness to the Division of Dockets Management (HFA-305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852.

Please cite the appropriate docket number as listed in table 1 of this document when submitting a written request. See the **SUPPLEMENTARY INFORMATION** section for electronic access to the summaries of safety and effectiveness.

FOR FURTHER INFORMATION CONTACT: Thinkh Nguyen, Center for Devices and Radiological Health (HFZ-402), Food and Drug Administration, 9200 Corporate Blvd., Rockville, MD 20850, 301-594-2186 ext. 152.

SUPPLEMENTARY INFORMATION:

ch066

NAL 1

I. Background

In the **Federal Register** of January 30, 1998 (63 FR 4571), FDA published a final rule that revised 21 CFR 814.44(d) and 814.45(d) to discontinue individual publication of PMA approvals and denials in the **Federal Register**. Instead, the agency now posts this information on the Internet on FDA's home page at <http://www.fda.gov>. FDA believes that this procedure expedites public notification of these actions because announcements can be placed on the Internet more quickly than they can be published in the **Federal Register**, and FDA believes that the Internet is accessible to more people than the **Federal Register**.

In accordance with section 515(d)(4) and (e)(2) of the Federal Food, Drug, and Cosmetic Act (the act) (21 U.S.C. 360e(d)(4) and (e)(2)), notification of an order approving, denying, or withdrawing approval of a PMA will continue to include a notice of opportunity to request review of the order under section 515(g) of the act. The 30 day period for requesting reconsideration of an FDA action under § 10.33(b) (21 CFR 10.33(b)) for notices announcing approval of a PMA begins on the day the notice is placed on the Internet. Section 10.33(b) provides that FDA may, for good cause, extend this 30 day period. Reconsideration of a denial or withdrawal of approval of a PMA may be sought only by the applicant; in these cases, the 30 day period will begin when the applicant is notified by FDA in writing of its decision.

The regulations provide that FDA publish a quarterly list of available safety and effectiveness summaries of PMA approvals and denials that were announced during that quarter. The following is a list of approved PMAs for which summaries of safety and effectiveness were placed on the Internet from October 1, 2005 through December 31, 2005. There were no denial actions

during this period. The list provides the manufacturer's name, the product's generic name or the trade name, and the approval date.

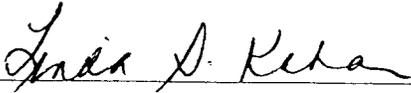
**Table 1.—List of Safety and Effectiveness Summaries for Approved PMAs
Made Available From October 1, 2005 through December 31, 2005**

PMA No./Docket No.	Applicant	Trade Name	Approval Date
P960040(S28)/2005M-0435	Guidant CRM Corp.	VENTAK PRIZM AVT AICD SYSTEM	March 27, 2003
P020045/2005M-0475	CryoCath Technologies, Inc.	7F FREEZOR CARDIAC CRYOABLATION CATHETER & CCT.2 CRYOCONSOLE SYSTEM	April 17, 2003
P040003/2005M-0473	InSightec—North America	EXABLATE 2000 SYSTEM	October 22, 2004
P030056/2005M-0478	Bayer Healthcare, LLC	ADVIA CENTAUR HCV READY PACK REAGENTS, ADVIA CENTAUR HCV QUALITY CONTROL MATE- RIALS	December 22, 2004
P980022(S11)/2005M-0454	Medtronic MiniMed	GUARDIAN RT CONTINUOUS GLU- COSE MONITORING SYSTEM	July 18, 2005
P020016/2005M-0399	Walter Lorenz Surgical, Inc.	TOTAL TEMPOMANDIBULAR JOINT REPLACEMENT SYSTEM	September 21, 2005
P040047/2005M-0477	Bioform Medical, Inc.	COAPTITE	November 10, 2005
P040042/2005M-0476	Irvine Biomedical, Inc.	THERAPY DUAL 8 CARDIAC ABLA- TION SYSTEM	November 18, 2005
P030054(S10)/2005M-0492	St. Jude Medical CRMD	EPIC & ATLAS + HF CRT-D SYS- TEMS	November 18, 2005
P040013/2005M-0474	Biomimetic Therapeutics, Inc.	GEM 21S (GROWTH-FACTOR EN- HANCED MATRIX)	November 18, 2005
P040045/2005M-0504	Vistakon, Division of Johnson & John- son Vision Care, Inc.	VISTAKON (SENOFILCON A) CON- TACT LENS, CLEAR AND VISI- BILITY TINTED WITH UV BLOCKER	December 20, 2005

II. Electronic Access

Persons with access to the Internet may obtain the documents at <http://www.fda.gov/cdrh/pmapage.html>.

Dated: 3/7/06
March 7, 2006.



Linda S. Kahan,
Deputy Director,
Center for Devices and Radiological Health.

[FR Doc. 06-????? Filed ??-??-06; 8:45 am]

BILLING CODE 4160-01-S

