



Food and Drug Administration  
9200 Corporate Boulevard  
Rockville MD 20850

Mr. Kris Shah  
Vice President, Product Development  
Baylis Medical Company, Inc.  
5160 Explorer Drive, Unit 33  
Mississauga, Ontario, Canada  
L4W 4T7

**MAY 03 2002**

Re: K020354

Trade/Device Name: Bayliss Pain Management Generator  
Model PMG - 115 (For Domestic Use)  
Model PMG - 230 (For International Use)

Regulation Number: 878.4400, 882.4400

Regulation Name: Electrosurgical cutting and coagulation device and accessories  
Radiofrequency lesion generator

Regulatory Class: II

Product Code: GEI, GXD

Dated: January 28, 2002

Received: February 4, 2002

Dear Mr. Shah:

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.

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), 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 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); 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.

This letter will allow you to begin marketing your device as described in your Section 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 21 CFR Part 809.10 for in vitro diagnostic devices), please contact the Office of Compliance at (301) 594-4659. 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 Part 807.97). Other general information on your responsibilities under the Act may be obtained from the Division of Small Manufacturers, International and Consumer 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*   
Celia M. Witten, Ph.D., M.D.  
Director  
Division of General, Restorative  
and Neurological Devices  
Office of Device Evaluation  
Center for Devices and  
Radiological Health

Enclosure

510(k) NUMBER (IF KNOWN): K020354

DEVICE NAME: Baylis Pain Management Generator  
Model: PMG – 115 (For Domestic Use)  
PMG – 230 (For International Use)

INDICATIONS FOR USE:

Baylis Pain Management Generator; Model PMG-115 (For Domestic Use) and Model PMG-230 (For International Use) is indicated for use to create lesions during neurological lesion procedures. The Baylis PMG is to be used with separately approved lesion/temperature probes such as Baylis Pain Management Probes and Oratec Spinecath™.

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use                        
(Per 21 CFR 801.109)

OR

Over-The-Counter-Use                     

Miriam C. Provost  
(Division Sign-Off)  
Division of General, Restorative  
and Neurological Devices

510(k) Number K020354