

DEPARTMENT OF HEALTH AND HUMAN SERVICES

PUBLIC HEALTH SERVICE
 FOOD AND DRUG ADMINISTRATION
 CDRH Medical Device Reporting
 P.O. Box 3002
 Rockville, MD 20847-3002

FORM APPROVED: OMB No. 0910-0437
 EXPIRES: 3/31/03

FOR FDA USE ONLY

MEDICAL DEVICE REPORTING BASELINE REPORT

PART 1

INSTRUCTIONS

Part 1 is a cover sheet for single or multiple copies of Part 2. The FDA Registration Number, (item 2.b.) and the Date of Baseline Report, (item 7.) must be provided on each attached Part 2. Return this form to the address listed above.

1. TYPE OF BASELINE REPORT
 Initial

 Annual Update
2. FIRM INFORMATION (Reporting Site)

a. Firm Name

b. FDA Registration Number (Reporting Site)

c. Street Address

d. City

e. State

f. ZIP Code

g. Country/Postal code (if not U.S.)

3. MANUFACTURER CONTACT INFORMATION

a. Name

b. Title

c. Street Address

d. City

e. State

f. ZIP Code

g. Telephone Number (Include area code and extension)

()

4. NUMBER OF "Baseline Reports - Part 2" attached**5. ARE YOU THE U.S. AGENT FOR A FOREIGN MANUFACTURER PER 21 CFR 803, 807?**
 YES

 NO
6. SIGNATURE**7. DATE**

_ / _ / _ _ _
 M M D D Y Y Y Y

Public reporting burden for this collection of information is estimated to average 1 hour per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden, to:

Center for Devices and Radiological Health
 Office of Surveillance and Biometrics
 Division of Surveillance Systems, RSMB, HFZ-533
 1350 Piccard Drive
 Rockville, MD 20850

An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number.

MEDICAL DEVICE REPORTING - BASELINE REPORT

 INITIAL

 ANNUAL UPDATE

PART 2

INSTRUCTIONS: Complete ONE copy of the following information for EACH device model. If model is not used, select another identifier from the device's labeling. Refer to baseline instructions, separate from this form, for detailed guidance.

FDA REGISTRATION NUMBER _____

DATE OF BASELINE _____ / _____ / _____

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1. MANUFACTURING SITE(S) FOR THIS DEVICE

a. FDA Registration Number (Manufacturing Site)

b. Firm Name (Manufacturing Site)

Note: If more space is required attach additional Part 2 pages, providing items 1.a. & 1.b. Include the FDA reporting site registration number (Part 1, item 2.b.), the date (Part 1, item 7.), device brand name (Part 2, item 2.), and model (Part 2, item 4.) for a cross reference on each page.

2. DEVICE BRAND NAME

3. DEVICE GENERIC NAME

4. DEVICE MODEL NUMBER

5. DEVICE CATALOG NUMBER

6. OTHER DEVICE IDENTIFIER

7. FDA PRODUCT CODE (Refer to FDA Classification Names booklet.)

8. MANUFACTURER'S DEVICE FAMILY NAME

9. RELATED DEVICE IDENTIFICATION If this device, or a substantially similar device, was previously distributed with a different device identification, provide the following:

a. Previous Device Identifier _____

b. Type of Identifier

- Model
 Catalog
 Other _____

c. Date of Prior Baseline Report

M M / D D / Y Y Y Y

10. BASIS FOR MARKETING

- a. 510 (k) _____ Yes, Number _____ No
 b. PMA _____ Yes, Number _____ No
 c. Preamendment ___ Yes No
 d. Transitional ___ Yes No
 e. 510 (k) Exempt ___ Yes No

11. DEVICE LIFE

- a. Shelf Life _____ months N/A

 Is shelf life labeled? (Go to 11.b. if N/A) Yes No

 b. Expected Life _____ months N/A Not Established/Indefinite

12. DATE DEVICE FIRST MARKETED

M M / D D / Y Y Y Y

13. DATE DEVICE CEASED BEING MARKETED (If applicable)

M M / D D / Y Y Y Y

14. IS THE DEVICE THE SUBJECT OF AN APPROVED POST MARKET (522) STUDY?

 Yes No

15. NUMBERS OF THIS DEVICE

Number

- a. Manufactured in the last 12 months _____
 b. Distributed in the last 12 months _____ Actual Estimated
 c. In current use _____ Actual Estimated

NOTE: Attach a copy of the method used to estimate the numbers in 15.b./15.c. OR if the method has been submitted in a previous baseline report complete item 16. below.

16. IF THE METHOD USED TO ESTIMATE THE NUMBERS IN 15.b. and/or 15.c. ABOVE HAS BEEN SUBMITTED IN A PREVIOUS BASELINE REPORT, COMPLETE THE FOLLOWING:

ID for device in prior baseline report
Date of previous baseline

Distribution Estimation Method

M M / D D / Y Y Y Y

Current Use Estimation Method

M M / D D / Y Y Y Y