

disregards income of a stepparent whose needs are not included in the assistance unit for the first 6-months of receipt of public assistance; excludes summer earnings of teens and interest income; lowers age of child for JOBS exemption to 6-months; raises asset limit to \$5,000 plus a vehicle of reasonable worth used primarily for self-sufficiency purposes; extends transitional Medicaid and child care benefits; eliminates 100-hour and required quarters of work rules, and (on a case-by-case basis) the 6-month time limit requirements in the AFDC-UP program; requires school conferences and regular school attendance; offers incentive payments to private employers who hire hard-to-place AFDC recipients; and allows non-custodial parents of AFDC children to participate in JOBS. Statewide, the demonstration requires immunizations of pre-school-age children.

Dated Received: 8/2/95.

Type: Combined AFDC/Medicaid.

Current Status: New.

Contact Person: Don Winstead, (904) 921-5567.

III. Requests for Copies of a Proposal

Requests for copies of this proposal should be directed to the Administration for Children and Families (ACF) at the address listed above. Questions concerning the content of the proposal should be directed to the State contact listed for the proposal.

(Catalog of Federal Domestic Assistance Program, No. 93562; Assistance Payments—Research).

Dated: August 11, 1995.

Howard Rolston,

Director, Office of Policy and Evaluation.

[FR Doc. 95-20293 Filed 8-15-95; 8:45 am]

BILLING CODE 4184-01-P

Food and Drug Administration

[Docket No. 95M-0119]

Chartex International plc; Premarket Approval of Femidom® Female Condom; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; correction.

SUMMARY: The Food and Drug Administration (FDA) is correcting a notice that appeared in the **Federal Register** of June 8, 1995 (60 FR 30310). The document announced the approval of the premarket approval application for the Femidom® Female Condom. The document was published with some errors. This document corrects those errors.

FOR FURTHER INFORMATION CONTACT: Marquita B. Steadman, Center for Devices and Radiological Health (HFZ-84), Food and Drug Administration, 2098 Gaither Rd., Rockville, MD 20850, 301-594-4765.

In FR Doc. 95-14059, appearing on page 30310 in the **Federal Register** of Thursday, June 8, 1995, the following corrections are made: On page 30310, in the second column, under the **SUMMARY** caption, in the fourth line, and under the **SUPPLEMENTARY INFORMATION** caption, in the second line, insert "Rhys, Bryant, U.S. representative for" before "Chartex International plc, London, U. K.,".

Dated: August 8, 1995.

Joseph A. Levitt,

Deputy Director for Regulations Policy, Center for Devices and Radiological Health.

[FR Doc. 95-20313 Filed 8-15-95; 8:45 am]

BILLING CODE 4160-01-F

Medical Devices; Mammography Facilities Education and Training; Notice of Public Workshops

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of public workshops.

SUMMARY: The Food and Drug Administration (FDA) (Office of Regulatory Affairs, Office of External Affairs, and Center for Devices and Radiological Health) is sponsoring five grassroots workshops on FDA requirements for compliance with the Mammography Quality Standards Act of 1992 (the MQSA). These workshops are designed to assist mammography facilities in complying with the regulations that went into effect on October 1, 1994.

DATES: The public workshops are scheduled as follows:

1. Thursday, August 17, 1995, 8 a.m. to 4:30 p.m., Dallas, TX.
2. Thursday, August 24, 1995, 8 a.m. to 4:30 p.m., Charlotte, NC.
3. Wednesday, September 6, 1995, 8 a.m. to 4:30 p.m., Fort Mitchell, KY.
4. Thursday, September 21, 1995, 8 a.m. to 4:30 p.m., San Juan, PR.
5. Thursday, September 28, 1995, 8 a.m. to 4:30 p.m., Los Angeles, CA.

ADDRESSES: The public workshops will be held at the following locations:

1. Dallas—Harvey Hotel, 400 North Olive, Dallas, TX.
2. Charlotte—New Charlotte Convention Center, 501 South College St., Charlotte, NC.
3. Fort Mitchell—Drawbridge Estates, 2477 Royal Dr., Fort Mitchell, KY.

4. Puerto Rico—Radisson Normandie Hotel, Avenida Munoz Rivera, Esquina Rosales, San Juan, PR.

5. Los Angeles—Continental Plaza, Los Angeles Airport, 9750 Airport Blvd., Los Angeles, CA. (PLEASE NOTE: Location changed since July 19, 1995, "Dear Colleague letter.")

FOR FURTHER INFORMATION CONTACT:

Regarding registration for the Dallas public workshop: Belinda Collins, Food and Drug Administration, Southwest Region, 7920 Elmbrook Rd., Dallas, TX 75247-4982, 214-655-8100, ext. 148 or FAX 214-655-8103.

Regarding registration for the Charlotte public workshop: Barbara Ward-Groves, Food and Drug Administration, Southeast Region, 60 Eighth St. SE., Atlanta, GA 30309, 404-347-4001, ext. 5256 or FAX 404-347-4349.

Regarding registration for the Fort Mitchell public workshop: Pat Wolfzorn, Food and Drug Administration, Mid-Atlantic Region, 1141 Central Pkwy., Cincinnati, OH 45202-1097, 513-684-3501, ext. 102 or FAX 513-684-2905.

Regarding registration for the San Juan public workshop: Nilda E. Villegas, Food and Drug Administration, Southeast Region, P. O. Box 5719, Puerta de Tierra Station, 809-729-6852 or FAX 809-729-6847.

Regarding registration for the Los Angeles public workshop: Mark Roh, Food and Drug Administration, Pacific Region, Oakland Federal Bldg., 1301 Clay St., suite 1180-N, Oakland, CA 94612-5217, 510-637-3980 or FAX 510-637-3977.

Those persons interested in attending a workshop should register by FAXing their name, firm name, address, and telephone number to the information contact person listed above for their region. There is no registration fee for these workshops, but advance registration is required. Interested parties are encouraged to register early because space is limited.

SUPPLEMENTARY INFORMATION: FDA will conduct training for mammography facilities designed to assist those facilities in complying with the requirements of the MQSA. Those requirements went into effect October 1, 1994. Emphasis will be placed on educational requirements, training, and providing assistance to small business in meeting the MQSA requirements. These meetings are being held, in part, as a response to the National