agency in developing a tentative final order. By this notice, FDA announces that it is treating these submissions, received after the administrative record has closed, as petitions to reopen the administrative record, and is granting the petitions by allowing new data and information contained therein to be included in the administrative record for OTC skin protectant drug products. This notice serves to inform interested persons of the existence of these data and information and their availability for review at the office of the Hearing Clerk, Food and Drug Administration. Comments on these data and information will not be accepted at this time. However, interested persons will have an opportunity to submit comments and additional new data and information at times to be specified in future **Federal Register** notices.

Dated: March 12, 1980.

William F. Randolph,

Acting Associate Commissioner for Regulatory Affairs.

[FR Doc. 80-8488 Filed 3-20-80; 8:45 am]

BILLING CODE 4110-03-M

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21 CFR Part 350

[Docket No. 78N-0064]

Antiperspirant Drug Products for Over-the-Counter Human Use

AGENCY: Food and Drug Administration.

ACTION: Acceptance of data and information into the administrative record.

SUMMARY: This notice advises that the Food and Drug Administration (FDA) reopened the administrative record for over-the counter (OTC) antiperspirant drug products to allow for consideration of data and information that had been filed with the Hearing Clerk, Food and Drug Administration, after the date that the administrative record officially closed.

ADDRESS: Data and information are on public file in the office of the Hearing Clerk (HFA-305), Food and Drug Administration, Rm. 4-62, 5600 Fishers Lane, Rockville, MD 20857, and may be seen between 9 a.m. and 4 p.m. Monday through Friday.

FOR FURTHER INFORMATION CONTACT: William E. Gilbertson, Bureau of Drugs (HFD-510), Food and Drug Administration, Department of Health, Education, and Welfare, 5600 Fishers Lane, Rockville, MD 20857, 301-443-4960.

SUPPLEMENTARY INFORMATION: The Food and Drug Administration has on numerous occasions received new data and information bearing on OTC drug panel reports and proposed monographs after the closing of the administrative record. Under § 330.10(a)(10)(i) (21 CFR 330.10(a)(10)(i)) the administrative record closes with respect to the submission of new data and information at the end of the comment period following publication of the panel report in the **Federal Register.** The comment period for OTC antiperspirant drug products closed on January 8, 1979, and the reply comment period closed on February 7, 1979. The procedural regulations for OTC drugs, § 330.10(a)(10)(ii) (21 CFR 330.10(a)(10)(ii)), provide that after the closing of the comment period following publication of the panel report, new data and information may be submitted for inclusion into the administrative record only through a petition to reopen the administrative record. In some cases, persons have not submitted such petitions; rather, they have submitted new data and information to the Hearing Clerk as comments on the panel report. In the interest of expediting the OTC drug review and because FDA wishes to consider all pertinent data and information that have been submitted to the Hearing Clerk, Food and Drug Administration, prior to the date of publication of this notice, the agency has concluded that the new data and information, whether or not properly filed, should be available to the agency in developing a tentative final order. By this notice, FDA announces that it is treating these submissions, received after the administrative record has closed, as petitions to reopen the administrative record, and is granting the petitions by allowing new data and information contained therein to be included in the Administrative record for OTC antiperspirant drug products. This notice serves to inform interested persons of the existence of these data and information and their availability for review at the office of the Hearing Clerk, Food and Drug Administration. Comments on these data and information will not be accepted at this time. However, interested persons will have an opportunity to submit comments and additional new data and information at times to be specified in future **Federal Register** notices.

Dated: March 12, 1980.

William F. Randolph,

Acting Associate Commissioner for Regulatory Affairs.

[FR Doc. 80-8470 Filed 3-20-80; 8:45 am]

BILLING CODE 4110-03-M

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21 CFR Part 352

[Docket No. 78N-0038]

Sunscreen Drug Products for Over-the-Counter Human Use

AGENCY: Food and Drug Administration.

ACTION: Acceptance of data and information into the administrative record.

SUMMARY: This notice advises that the Food and Drug Administration (FDA) reopened the administrative record for over-the-counter (OTC) sunscreen drug products to allow for consideration of data and information that had been filed with the Hearing Clerk, Food and Drug Administration, after the date that the administrative record officially closed.

ADDRESS: Data and information are on public file in the office of the Hearing Clerk (HFA-305), Food and Drug Administration, Rm. 4-62, 5600 Fishers Lane, Rockville, MD 20857, and may be seen between 9 a.m. and 4 p.m., Monday through Friday.

FOR FURTHER INFORMATION CONTACT: William E. Gilbertson, Bureau of Drugs (HFD-510), Food and Drug Administration, Department of Health, Education, and Welfare, 5600 Fishers Lane, Rockville, MD 20857, 301-443-4960.

SUPPLEMENTARY INFORMATION: The Food and Drug Administration has on numerous occasions received new data and information bearing on OTC drug panel reports and proposed monographs after the closing of the administrative record. Under § 330.10(a)(10)(i) (21 CFR 330.10(a)(10)(i)) the administrative record closes with respect to the submission of new data and information at the end of the comment period following publication of the panel report in the **Federal Register.** The comment period for OTC sunscreen drug products closed on November 24, 1978, and the reply comment period closed on December 26, 1978. The procedural regulations for OTC drugs, § 330.10(a)(10)(ii) (21 CFR 330.10(a)(10)(ii)), provide that after the closing of the comment period following publication of the panel report, new data and information may be submitted for inclusion into the administrative record only through a petition to reopen the administrative record. In some cases, persons have not submitted such petitions; rather, they have submitted new data and information to the Hearing Clerk as comments on the panel report. In the interest of expediting the OTC drug review and because FDA wishes to consider all pertinent data and information that have been

submitted to the Hearing Clerk, Food and Durg Administration, prior to the date of publication of this notice, the agency has concluded that the new data and information, whether or not properly filed, should be available to the agency in developing a tentative final order. By this notice, FDA announces that it is treating these submissions, received after the administrative record has closed, as petitions to reopen the administrative record, and is granting the petitions by allowing new data and information contained therein to be included in the administrative record for OTC sunscreen drug products. This notice serves to inform interested persons of the existence of these data and information and their availability for review at the office of the Hearing Clerk, Food and Drug Administration. Comments on these data and information will not be accepted at this time. However, interested persons will have an opportunity to submit comments and additional new data and information at times to be specified in future **Federal Register** notices.

Dated: March 12, 1980.

William F. Randolph,

Acting Associate Commissioner for Regulatory Affairs.

[FR Doc. 80-8489 Filed 3-20-80; 8:45 am]

BILLING CODE 4110-03-M

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21 CFR Part 358

[Docket No. 78N-0065]

Skin Bleaching Drug Products for Over-the-Counter Human Use

AGENCY: Food and Drug Administration.

ACTION: Acceptance of data and information into the administrative record.

SUMMARY: This notice advises that the Food and Drug Administration (FDA) reopened the administrative record for over-the-counter (OTC) skin bleaching drug products to allow for consideration of data and information that had been filed with the Hearing Clerk, Food and Drug Administration, after the date that the administrative record officially closed.

ADDRESS: Data and information are on public file in the office of the Hearing Clerk (HFA-305), Food and Drug Administration, Rm. 4-62, 5600 Fishers Lane, Rockville, MD 20857, and may be seen between 9 a.m. and 4 p.m., Monday through Friday.

FOR FURTHER INFORMATION CONTACT: William E. Gilbertson, Bureau of Drugs (HFD-510), Food and Drug Administration, Department of Health, Education, and Welfare, 5600 Fishers Lane, Rockville, MD 20857, 301-443-4960.

SUPPLEMENTARY INFORMATION: The Food and Drug Administration has on numerous occasions received new data and information bearing on OTC drug panel reports and proposed monographs after the closing of the administrative record. Under § 330.10(a)(10)(i) (21 CFR 330.10(a)(10)(i)) the administrative record closes with respect to the submission of new data and information at the end of the comment period following publication of the panel report in the **Federal Register.** The comment period for OTC skin bleaching drug products closed on February 1, 1979, and the reply comment period closed on March 5, 1979. The procedural regulations for OTC drugs, § 330.10(a)(10)(ii) (21 CFR 330.10(a)(10)(ii)), provide that after the closing of the comment period following publication of the panel report, new data and information may be submitted for inclusion into the administrative record only through a petition to reopen the administrative record. In some cases, persons have not submitted such petitions; rather they have submitted new data and information to the Hearing Clerk as comments on the panel report. In the interest of expediting the OTC drug review and because FDA wishes to consider all pertinent data and information that have been submitted to the Hearing Clerk, Food and Drug Administration, prior to the date of publication of this notice, the agency has concluded that the new data and information, whether or not properly filed, should be available to the agency in developing a tentative final order. By this notice, FDA announces that it is treating these submissions, received after the administrative record has closed, as petitions to reopen the administrative record, and is granting the petitions by allowing new data and information contained therein to be included in the administrative record for OTC skin bleaching drug products. This notice serves to inform interested persons of the existence of these data and information and their availability for review at the office of the Hearing Clerk, Food and Drug Administration. Comments on these data and information will not be accepted at this time. However, interested persons will have an opportunity to submit comments and additional new data and information at times to be specified in future **Federal Register** notices.

Dated: March 12, 1980.

William F. Randolph,

Acting Associate Commissioner for Regulatory Affairs.

[FR Doc. 80-8505 Filed 3-20-80; 8:45 am]

BILLING CODE 4110-03-M

DEPARTMENT OF HOUSING AND URBAN DEVELOPMENT

Office of the Secretary

24 CFR Part 570

[Docket No. R-80-782]

Community Development Block Grants; Small Cities Program; Transmittal of Proposed Rule to Congress

AGENCY: Department of Housing and Urban Development.

ACTION: Notice of transmittal of proposed rule to Congress under Section 7(o) of the Department of HUD Act.

SUMMARY: Recently enacted legislation authorizes Congress to review certain HUD rules for fifteen (15) calendar days of continuous session of Congress prior to each such rule's publication in the **Federal Register.** This Notice lists and summarizes for public information a proposed rule which the Secretary is submitting to Congress for such review.

FOR FURTHER INFORMATION CONTACT: Burton Bloomberg, Director, Office of Regulations, Office of General Counsel, 451 7th Street SW., Washington, D.C. 20410, (202) 755-6207.

SUPPLEMENTARY INFORMATION: Concurrently with issuance of this Notice, the Secretary is forwarding to the Chairmen and Ranking Minority Members of both the Senate Banking, Housing and Urban Affairs Committee and the House Banking, Finance and Urban Affairs Committee the following proposed rulemaking document:

24 CFR Part 570, Subpart F—Community Development Block Grants—Small Cities Program (FY 1981)

This proposed rule would revise the regulations for the Small Cities Program for Community Development Block Grants. Most of the changes would clarify existing policies and procedures, but some substantive changes with respect to procedures and program standard are proposed. The important changes are:

1. Elimination of the housing needs factors as part of the selection system;
2. A new method of computing the poverty needs factors;
3. Replacing criteria dealing with State policies and a community's