time for submission to the objective review group. (The applicants must request a legibly dated U.S. Postal Service postmark or obtain a legibly dated receipt from a commercial carrier or the U.S. Postal Service. Private metered postmarks shall not be acceptable as proof of timely mailing.)

2. Late Applicants: Applications that do not meet the criteria in 1.(a) or 1.(b) above are considered late applications. Late applications will not be considered in the current competition and will be returned to the applicants.

Where to Obtain Additional Information

To receive additional written information call (404) 332–4561. You will be asked to leave your name, address, and telephone number and will need to refer to Announcement 528. You will be receive a complete program description, information on application procedures, and application forms.

If you have questions after reviewing the contents of all the documents, business management technical assistance may be obtained from Oppie M. Byrd, Grants Management Specialist, Grants Management Branch, Procurement and Grants Office, Centers for Disease Control and Prevention (CDC), 255 East Paces Ferry Road, NE., Room 300, Mailstop E-13, Atlanta, GA 30305, telephone (404) 842-6546. Programmatic technical assistance may be obtained from Marie Haring Sweeney, Ph.D., National Institute for Occupational Safety and Health, Division of Surveillance, Hazard Evaluation and Field Studies, Centers for Disease Control and Prevention (CDC), Mailstop R-13, Robert A. Taft Laboratories, 4676 Columbia Parkway, Cincinnati, OH 45226-1049, telephone (513) 841-4207.

Potential applicants may obtain a copy of Healthy People 2000 (Full Report, Stock No. 017–001–00474–0) or Healthy People 2000 (Summary Report, Stock No. 017–001–00473–1) referenced in the INTRODUCTION through the Superintendent of Documents, Government Printing Office, Washington, DC 20402–9325, telephone (202) 512–1800.

Copies of A Framework for Assessing the Effectiveness of Disease and Injury Prevention (CDC, Morbidity and Mortality Weekly Report, March 27, 1992, Volume 41, Number RR–3, pages 5–11) may be obtained by calling (404) 488–4334.

Dated: April 21, 1995.

Diane D. Porter,

Acting Director, National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention.

[FR Doc. 95-10454 Filed 4-27-95; 8:45 am]

BILLING CODE 4163-19-P

Food and Drug Administration

[Docket No. 95C-0091]

GNT Gesellshaft für Nahrungsmitteltechnologie mbH; Filing of Color Additive Petition

AGENCY: Food and Drug Administration,

HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that GNT Gesellshaft für Nahrungsmitteltechnologie mbH has filed a petition proposing that the color additive regulations be amended to provide for the safe use of dried fruit juice color additive, dried vegetable juice color additive, and vegetable juice color additive prepared by water infusion of the dried vegetable.

FOR FURTHER INFORMATION CONTACT: Aydin Örstan, Center for Food Safety and Applied Nutrition (HFS–217), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202–418–3076.

SUPPLEMENTARY INFORMATION: Under the Federal Food, Drug, and Cosmetic Act (sec. 721(b)(5) (21 U.S.C. 379e(b)(5))), notice is given that a color additive petition (CAP 5C0245) has been filed by GNT Gesellshaft für Nahrungsmitteltechnologie mbH, c/o Burditt & Radzius, Chtd., 333 West Wacker Dr., suite 2600, Chicago, IL 60606-1218. The petition proposes to amend the color additive regulations in § 73.250 Fruit juice (21 CFR 73.250) to provide for the safe use of dried fruit juice color additive and in § 73.260 Vegetable juice (21 CFR 73.260) to provide for the safe use of dried vegetable juice color additive, and vegetable juice color additive prepared by water infusion of the dried vegetable.

The agency has determined under 21 CFR 25.24(a)(9) that this action is of a type that does not individually or cumulatively have a significant effect on the human environment. Therefore, neither an environmental assessment nor an environmental impact statement is required.

Dated: April 20, 1995.

Alan M. Rulis,

Acting Director, Office of Premarket Approval, Center for Food Safety and Applied Nutrition.

[FR Doc. 95–10539 Filed 4–27–95; 8:45 am]

BILLING CODE 4160-01-F

[Docket No. 93N-0418]

United Blood Services Blood Systems, Inc.; Revocation of U.S. License No. 0183–020

AGENCY: Food and Drug Administration,

HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the revocation of the establishment license (U.S. License No. 0183-020) and product licenses issued to United Blood Services Blood Systems, Inc. (BSI), for the manufacture of Whole Blood (ACD, CPD, CPDA-1), Red Blood Cells, Red Blood Cells Leukocytes Removed, Plasma, Fresh Frozen Plasma, Cryoprecipitated AHF, Platelets, Platelets Pheresis, and Source Leukocytes. BSI has numerous locations throughout the United States; the licenses have been revoked only at the BSI location at Texarkana, TX. In a letter to FDA dated June 28, 1993, BSI voluntarily requested the revocation of its establishment and product licenses and waived its opportunity for hearing. **DATES:** The revocation of the establishment license (U.S. License No.

establishment license (U.S. License No. 0183–020) and the product licenses became effective July 23, 1993.

FOR FURTHER INFORMATION CONTACT: Jean M. Olson, Center for Biologics Evaluation and Research (HFM–635), Food and Drug Administration, 1401 Rockville Pike, Rockville, MD 20852–1448, 301–594–3074.

SUPPLEMENTARY INFORMATION: FDA is announcing the revocation of the establishment license (U.S. License No. 0183-020) and the product licenses issued to BSI, 1321 College Dr., Texarkana, TX 75503, for the manufacture of Whole Blood (ACD, CPD, CPDA-1), Red Blood Cells, Red Blood Cells Leukocytes Removed, Plasma, Fresh Frozen Plasma, Cryoprecipitated AHF, Platelets, Platelets Pheresis, and Source Leukocytes. The current mailing address is United Blood Services Blood Systems, Inc., c/o Blood Systems, Inc., 6210 East Oak St., P.O. Box 1867, Scottsdale, AZ 85252. BSI has numerous locations throughout the United States. The licenses were revoked for the Texarkana, TX, location only.