ful activity. If a person's earnings from work activity do not show whether he or she has done or has not done substantial gainful activity under paragraph (b)(1) or (b)(2) of this section, we will consider other information in addition to earnings. We will look at the circumstances surrounding the work activity (see §§ 404.1532 and 404.1533 and paragraph (a) of this section) together with the medical evidence of the impairment or impairments and other factors (see § 404.1502 about evaluating impairments). We will ordinarily consider the earnings of a person working in a sheltered workshop or comparable facility especially set up for severely impaired persons as showing he or she is not doing substantial gainful activity if—

(i) The work activity is restricted by his or her impairment; and

(ii) The average monthly earnings for the particular calendar year are not more than the dollar limits contained in paragraphs (b)(1)(i) through (b)(1)(v) of this section.

(c) [Reserved]

(d) [Reserved]

(e) *Persons who are self-employed.*

* * *

* * * * *

2. Section 416.934 is amended by revising paragraph (b), revoking and reserving paragraphs (c) and (d), and revising the heading of paragraph (e) to read as follows:

§ 416.934 Evaluation of earnings from work.

* * * * *

(b) *Earnings guidelines.* We will use the following guidelines in paragraphs (b) (1), (2), and (3) of this section to determine whether a person's earnings show the ability to do substantial gainful activity.

(1) *Earnings that will ordinarily show that a person has done substantial gainful activity.* We will ordinarily consider that a person's earnings from work activity show that he or she has done substantial gainful activity if—

(i) Earnings averaged more than $200 a month in calendar years prior to 1976;

(ii) Earnings averaged more than $230 a month in calendar year 1976;

(iii) Earnings average more than $240 a month in calendar year 1977;

(iv) Earnings averaged more than $260 a month in calendar year 1978; or

(v) Earnings averaged more than $280 a month in calendar years after 1978; and

(vi) There is no evidence that the person was unable to do substantial gainful activity under the rules in §§ 416.932 and 416.933 and paragraph (a) of this section.

(2) *Earnings that will ordinarily show that a person has not done substantial gainful activity.* We will ordinarily consider that a person's earnings from work activity as an employee show that he or she has not done substantial gainful activity if—

(i) Earnings averaged less than $130 a month in calendar years prior to 1976;

(ii) Earnings averaged less than $150 a month in calendar year 1976;

(iii) Earnings averaged less than $160 a month in calendar year 1977;

(iv) Earnings averaged less than $170 a month in calendar year 1978; or

(v) Earnings averaged less than $180 a month in calendar years after 1978; and

(vi) There is no evidence that the person was able to do substantial gainful activity under the rules in §§ 416.932 and 416.933 and paragraph (a) of this section.

(3) *Earnings that are not high or low enough to show whether a person has done or has not done substantial gainful activity.* If a person's earnings from work activity do not show whether he or she has done or has not done substantial gainful activity under paragraph (b)(1) or (b)(2) of this section, we will consider other information in addition to earnings. We will look at the circumstances surrounding the work activity (see §§ 416.932 and 416.933 and paragraph (a) of this section) together with the medical evidence of the impairment or impairments and other factors (see § 416.902 about evaluating impairments). We will ordinarily consider the earnings of a person working in a sheltered workshop or comparable facility especially set up for severely impaired persons as showing he or she is not doing substantial gainful activity if—

(i) The work activity is restricted by his or her impairment; and

(ii) The average monthly earnings for the particular calendar year are not more than the dollar limits contained in paragraphs (b)(1)(i) through (b)(1)(v) of this section.

(c) [Reserved]

(d) [Reserved]

(e) *Persons who are self-employed.*

* * *

* * * * *

[FR Doc. 79-8087 Filed 3-22-79; 8:45 am]

[4510-27-M]

Title 20—Employee Benefits

CHAPTER VI—EMPLOYMENT STANDARDS ADMINISTRATION, DEPARTMENT OF LABOR

SUBCHAPTER A—LONGSHOREMEN'S AND HARBOR WORKERS' COMPENSATION ACT AND RELATED STATUTES

PART 702—ADMINISTRATION AND PROCEDURES

OFFICE OF WORKERS' COMPENSATION PROGRAMS; LONGSHOREMEN'S AND HARBOR WORKERS' COMPENSATION

Transfer of Districts

AGENCY: Employment Standards Administration, Labor.

ACTION: Final rule.

SUMMARY: The area and states formerly served by the Longshoremen's and Harbor Workers' Compensation Act District Office in Cleveland, Ohio are transferred to the Chicago District Office. This action is taken to better serve the public and provide improved administration of the Longshoremen's and Harbor Workers' Compensation Act.

DATE: This amendment shall be effective April 2, 1979.

FOR FURTHER INFORMATION CONTACT:

John E. Stocker, Associate Director for Longshore and Harbor Workers' Compensation, U.S. Department of Labor, Washington, D.C. 20210 (202) 523-8721.

SUPPLEMENTARY INFORMATION: This document changes compensation districts under § 702.101 of Part 702, Title 20 of the Code of Federal Regulations, establishing compensation districts pursuant to section 39(b) of the Longshoremen's and Harbor Workers' Compensation Act and its extensions.

The boundaries of existing district 10 are revised as follows: the States of Indiana, Michigan and Ohio are transferred from the heretofore existing district 9, with headquarters in Cleveland, Ohio, to the jurisdiction of district 10. All functions performed in district 9 are hereby transferred to district 10. Chicago remains the headquarters city of district 10. As of this date, district 10 is comprised of the States of Illinois, Indiana, Michigan, Minnesota, Ohio and Wisconsin.

The regulations at 20 CFR 704.101, 704.201, 704.301 and 704.401 are unchanged by this amendment.

Accordingly, Part 702 is amended as follows:

Section 702.101 is amended to show elimination of district 9 from the list as follows:

§ 702.101 Establishment of compensation districts.

Pursuant to section 39(b) of the Longshoremen's and Harbor Workers' Compensation Act, the following compensation districts have been established:

* * * * *

District No. 8. Comprises the States of Texas, Oklahoma, and New Mexico, with headquarters in Houston, Texas. District No. 10. Comprises the States of Illinois, Indiana, Michigan, Minnesota, Ohio and Wisconsin, with headquarters at Chicago, Illinois.

* * * * *

Signed at Washington, D.C. the 14th day of March 1979.

DONALD ELISBURG,
Assistant Secretary of Labor.

[FR Doc. 79-8990 Filed 3-22-79; 8:45 am]

[4110-03-M]

Title 21—Food and Drugs

CHAPTER I—FOOD AND DRUG ADMINISTRATION, DEPARTMENT OF HEALTH, EDUCATION, AND WELFARE

SUBCHAPTER F—BIOLOGICS

ZIP CODE CHANGE

AGENCY: Food and Drug Administration.

ACTION: Editorial amendments.

SUMMARY: The agency is amending 21 CFR Subchapter F (Parts 600 through 680) to change the zip code number.

EFFECTIVE DATE: March 23, 1979.

FOR FURTHER INFORMATION CONTACT:

Albert Rothschild, Bureau of Biologics (HFB-620), Food and Drug Administration, Department of Health, Education, and Welfare, 8800 Rockville Pike, Bethesda, MD 20205, 301-443-1306.

SUPPLEMENTARY INFORMATION: The zip code for the Bureau of Biologics has been changed. Because the zip code appears throughout the biologics regulations in 21 CFR Subchapter F, the agency is amending the Code of Federal Regulations in 21 CFR Parts 600 through 680 to reflect this change.

Accordingly, Title 21, Chapter I, of the Code of Federal Regulations is amended in Subchapter F, Parts 600 through 680 by changing the zip code number "20014" wherever it appears, to read "20205."

Dated: March 16, 1979.

WILLIAM F. RANDOLPH,
Acting Associate Commissioner for Regulatory Affairs.

[FR Doc. 79-8618 Filed 3-22-79; 8:45 am]

[4110-03-M]

CHAPTER I—FOOD AND DRUG ADMINISTRATION, DEPARTMENT OF HEALTH, EDUCATION, AND WELFARE

[Docket No. 76N-0400]

NONCLINICAL LABORATORY STUDIES

Good Laboratory Practice Regulations; Correction

AGENCY: Food and Drug Administration.

ACTION: Correction.

SUMMARY: In FR Doc. 78-35272 appearing at page 59986 in the FEDERAL REGISTER of Friday, December 22, 1978, the corrections set forth below are made.

FOR FURTHER INFORMATION CONTACT:

Paul D. Lepore, Bureau of Veterinary Medicine (HFV-102), Food and Drug Administration, Department of Health, Education, and Welfare, 5600 Fishers Lane, Rockville, Md., 301-443-4313.

CORRECTIONS

1. On page 59996, paragraph 71 of the preamble is corrected in the 3rd line by changing "responses to" read "responses of."

2. On page 59998, in the right column, paragraph 91 of the preamble is corrected in the 9th and 10th lines by changing "Although the Commissioner is deleting the requirement" to read "Accordingly, the Commissioner is requiring."

3. On page 60008, paragraph 196 of the preamble is corrected by changing "§ 58.185(a)(12)" to read "§ 58.185 (a)(10)."

4. On page 60009, paragraph 207 of the preamble is corrected in the 17th line by changing "marketed-sense" to read "mark-sense."

5. In Part 58, § 58.33 *Study director* is corrected in paragraph (b) by changing "responses to" to read "responses of."

6. In Part 601, § 601.2 *Application for establishment and product licenses; procedures for filing* is corrected in paragraph (a) by adding before the last sentence the sentence "The applicant shall also include an environmental impact analysis report analyzing the environmental impact of the manufacturing process and the ultimate use or consumption of the biological product pursuant to § 25.1 of this chapter."

7. On page 60024, amendment 18b is corrected to read as follows:

b. In § 1010.5, by revising the introductory text of paragraph (c) and by adding new paragraph (c)(13) to read as follows:

§ 1010.5 Exemptions for products intended for United States Government use.

* * * * *

(c) *Application for exemption.* An application for exemption, or for amendment or extension thereof, shall be submitted in quintuplicate to the Hearing Clerk, Food and Drug Administration, Rm. 4-65, 5600 Fishers Lane, Rockville, MD 20857. For an exemption pursuant to the criteria prescribed in paragraph (a)(1) of this section, the application shall include the information prescribed in paragraph (c)(1) through (13) of this section. For an exemption pursuant to the criteria prescribed in paragraph (a)(2) of this section, the application shall include the information prescribed in paragraph (c)(3) through (13) of this section. An application for exemption, or for amendment or extension thereof, and correspondence relating to such application shall be made available for public disclosure in the office of the Hearing Clerk, except for confidential or proprietary information submitted in accordance with Part 4 of this chapter. Information classified for reasons of national security shall not be included in the application. Except as indicated above, the application for exemption shall include the following:

* * * * *

(13) With respect to each nonclinical study contained in the application, either a statement that the study was conducted in compliance with the good laboratory practice regulations set forth in Part 58 of this chapter, or, if the study was not conducted in compliance with such regulations, a statement that describes in detail all differences between the practices used in the study and those required in the regulations.

* * * * *

FOR FURTHER INFORMATION CONTACT:

Paul D. Lepore, Bureau of Veterinary Medicine (HFV-102), Food and Drug Administration, Department of

Health, Education, and Welfare, 5600 Fishers Lane, Rockville, MD., 301-443-4313.

Dated: March 19, 1979.

WILLIAM F. RANDOLPH,
Acting Associate Commissioner for Regulatory Affairs.

[FR Doc. 79-9094 Filed 3-22-79; 8:45 am]

[4110-03-M]

SUBCHAPTER A—GENERAL

SUBCHAPTER F—BIOLOGICS

EDITORIAL AMENDMENTS

AGENCY: Food and Drug Administration.

ACTION: Correction.

SUMMARY: The Food and Drug Administration is making three editorial amendments to its regulations contained in Parts 74, 80, and 640.

EFFECTIVE DATE: March 23, 1979.

FOR FURTHER INFORMATION CONTACT:

John Richards, Federal Register Writer (HFC-11), Food and Drug Administration, Department of Health, Education, and Welfare, 5600 Fishers Lane, Rockville, MD 20857, 301-443-2994.

SUPPLEMENTARY INFORMATION: The agency is amending the Code of Federal Regulations in Parts 74, 80, and 640 as follows:

1. In Part 74, § 74.705 *FD&C Yellow No. 5* is amended in paragraph (b) in its list of specifications by changing in the 3d item "0.2 percent" to read "0.1 percent."

2. In Part 80, § 80.21 *Request for certification* is amended in paragraph (j)(1) by changing "Batch weighs" to read "Batch weights."

3. In Part 640, § 640.33 *Testing the blood* is amended in paragraph (a) by changing "§ 640.5(a), (b), and (e)" to read "§ 640.5(a), (b), and (c)."

Effective date: March 23, 1979.

Dated: March 19, 1979.

WILLIAM F. RANDOLPH,
Acting Associate Commissioner for Regulatory Affairs.

[FR Doc. 79-8997 Filed 3-22-79; 8:45 am]

[6560-01-M]

[FRL 1080-8; FAP 6H5126/R45]

SUBCHAPTER B—FOOD AND FOOD PRODUCTS

PART 193—TOLERANCES FOR PESTICIDES IN FOOD ADMINISTERED BY THE ENVIRONMENTAL PROTECTION AGENCY

SUBCHAPTER E—ANIMAL FEEDS, DRUGS, AND RELATED PRODUCTS

PART 561—TOLERANCES FOR PESTICIDES IN ANIMAL FEEDS ADMINISTERED BY THE ENVIRONMENTAL PROTECTION AGENCY

Glyphosate

AGENCY: Office of Pesticide Programs, Environmental Protection Agency (EPA).

ACTION: Final rule.

SUMMARY: This rule establishes tolerances for residues of the herbicide glyphosate in sugarcane molasses. The regulations were requested by Monsanto Co. This rule establishes maximum permissible levels for residues of glyphosate in sugarcane molasses.

EFFECTIVE DATE: March 23, 1979.

FOR FURTHER INFORMATION CONTACT:

Mr. Robert Taylor, Product Manager (PM) 25, Registration Division (TS-767), Office of Pesticide Programs, EPA, 401 M Street, SW, Washington, DC 20460 (202/755-7013).

SUPPLEMENTARY INFORMATION: On December 5, 1978, the EPA published in the FEDERAL REGISTER (43 FR 57005) a notice of proposed rulemaking to amend 21 CFR 193.235 and 561.253 by establishing regulations permitting combined residues of the herbicide glyphosate (*N*-(phosphonomethyl)glycine) and its metabolite aminomethylphosphonic acid in or on sugarcane molasses resulting from the application of the herbicide to growing sugarcane with a tolerance limitation of 2 parts per million (ppm). This notice was published in connection with a petition (FAP 6H5126) submitted by Monsanto Agricultural Products Co., 800 N. Lindbergh Blvd., St. Louis, MO 63166. (A related document establishing a tolerance for residues of glyphosate on sugarcane appears elsewhere in today's FEDERAL REGISTER.) No comments or requests for referral to an advisory committee were received in response to this notice of proposed rulemaking.

For purposes of clarification, the Agency has determined that the regulations should specify that the residues result from the herbicide application of the isopropylamine salt of glyphosate.

The data submitted in the petition and other relevant material have been evaluated, and it is concluded that the pesticide may be safely used in the prescribed manner when such use is in accordance with the label and labeling registered pursuant to the Federal Insecticide, Fungicide, and Rodenticide Act (FIFRA), as amended in 1972, 1975, and 1978 (92 Stat. 819; 7 U.S.C. 136). Therefore, the regulations are being established.

Any person adversely affected by this regulation may, on or before April 23, 1979, file written objections with the Hearing Clerk, Environmental Protection Agency, Rm. M-3708, 401 M St., SW, Washington, DC 20460. Such objections should be submitted and specify the provisions of the regulation deemed to be objectionable and the grounds for the objections. If a hearing is requested, the objections must state the issues for the hearing. A hearing will be granted if the objections are supported by grounds legally sufficient to justify the relief sought.

Effective on March 23, 1979, 21 CFR 193 and 561 are amended as set forth below.

(Section 409(c)(1) of the Federal Food, Drug, and Cosmetic Act [21 U.S.C. 348(c)(1)].)

Dated: February 24, 1979.

EDWIN L. JOHNSON,
Deputy Assistant Administrator for Pesticide Programs.

1. Part 193, Subpart A, § 193.235, is amended by revising paragraph (e) to read as follows:

§ 193.235 Glyphosate.

* * * * *

(e) Tolerances are established for combined residues of glyphosate (*N*-(phosphonomethyl)glycine) and its metabolite aminomethylphosphonic acid in the following processed foods when present therein as a result of herbicide application of the isopropylamine salt of glyphosate in palm tree culture and to growing sugarcane:

2 parts per million in sugarcane molasses.

0.1 part per million in palm oil.

2. Part 561, § 561.253, is amended by revising paragraph (f) to read as follows:

§ 561.253 Glyphosate.

* * * * *

(f) Tolerances are established for combined residues of glyphosate (*N*-(phosphonomethyl)glycine) and its metabolite aminomethylphosphonic acid in or on the following processed feeds when present therein as a result