may be handled during the period September 7, 1975, through September 13, 1975, is hereby fixed at 225,000 cartons.

(2) As used in this section, "handled", and "carton(s)" have the same meaning as when used in the said amended marketing agreement and order.

(Secs. 1–19, 48 Stat. 31, as amended (7 U.S.C. 601–674))

Dated: September 3, 1975.

CHARLES R. BRADER,
Deputy Director, Fruit and Vegetable Division, Agricultural Marketing Service.

[FR Doc.75-23842 Filed 9-4-75;11:22 am]

CHAPTER XIV—COMMODITY CREDIT CORPORATION, DEPARTMENT OF AGRICULTURE

[CCC Grain Price Support Regulations, 1975 Crop Peanut Farm-Stored Loan and Purchase Supplement]

PART 1421—GRAINS AND SIMILARLY HANDLED COMMODITIES

Loan and Purchase Rates

On January 27, 1975, notice of proposed rulemaking regarding loan and purchase rates for 1975 crop peanuts and operating provisions to carry out the 1975 crop peanut loan and purchase program was published in the FEDERAL REGISTER, 40 FR 4019.

Six responses were received from individual producers and other interested parties. None of the written comments, suggestions, or objections received, pertained to the aspects of the loan and purchase program covered by this subpart. The regulations contained in 7 CFR, §§ 1421.291 through 1421.295 are revised to read as follows, effective as to the 1975 crop of peanuts. The material previously appearing in these sections remains in full force and effect as to the crops to which it was applicable.

Subpart—1975 Crop Farm Stored Peanut Loan and Purchase Program

Sec.
1421.291 Purpose.
1421.292 Availability.
1421.293 Maturity of loans.
1421.294 Loan and purchase rates.
1421.295 Eligible peanuts.

AUTHORITY: Secs. 4 and 5, 62 Stat. 1070, as amended (15 U.S.C. 714 (b) and (c)); Secs. 101, 401, 403 and 405, 63 Stat. 1051, as amended (7 U.S.C. 1441, 1421, 423, 425).

Subpart—1975 Crop Farm Stored Peanut Loan and Purchase Program

§ 1421.291 Purpose.

This supplement contains program provisions which, together with the applicable provisions of the General Regulations Governing Price Support for the 1970 and Subsequent Crops of Grains and Similarly Handled Commodities (35 FR 7363 and 7761, 7 CFR 1421.1–29), as amended, and the provisions of the 1970 and Subsequent Crops Peanut Farm-Stored Loan and Purchase Supplement (35 FR 12706, 7 CFR 1421.280–289), as amended, (hereinafter referred to as "the continuing supplement"), which contain regulations of a general nature with respect to loan and purchase operations, apply to farm-stored loans and purchases for the 1975 crop of peanuts.

§ 1421.292 Availability.

(a) *Farm-stored loans.* Producers must request a loan on 1975 crop eligible peanuts on or before March 31, 1976.

(b) *Purchases.* Producers desiring to offer for purchase, eligible peanuts not under loan must execute and deliver to the appropriate county ASCS office, on or before April 30, 1976, a Purchase Agreement (Form CCC–614) indicating the approximate quantity of 1975 crop peanuts he may sell to CCC.

§ 1421.293 Maturity of loans.

Unless demand is made earlier, farm-stored loans on farmers' stock peanuts will mature on April 30, 1976.

§ 1421.294 Loan and purchase rates.

(a) *Loan rate.* Subject to the discounts specified in paragraph (b) of this section, the loan rates for farmers' stock peanuts placed under farm-stored loan shall be the following rates by types per ton:

Type:	Dollars per ton
Virginia	394
Runner	399
Southeast Spanish	385
Southwest Spanish	381
Valencia (suitable for cleaning and roasting in southwest)[1]	394

(b) *Location adjustment to support prices.* The loan rates specified in paragraph (a) of this section shall be subject to the following discounts for farmers' stock peanuts placed under a farm-stored loan in the States specified where peanuts are not customarily shelled or crushed:

State:	Dollars per ton
Arizona	25
Arkansas	10
California	33
Louisiana	7
Mississippi	10
Missouri	10
Tennessee	25

(c) *Settlement values.* The support prices, premiums, and discounts for use in computing the settlement value, under § 1421.289(b)(2) of the continuing supplement, of peanuts acquired by CCC under loan or purchase shall be those specified in § 1446.12 of the 1975 crop peanut warehouse storage loan supplement (39 FR 26715, 7 CFR 1446.12), including the location adjustments specified therein for peanuts delivered to CCC in States where peanuts are not customarily shelled or crushed.

§ 1421.295 Eligible peanuts.

In addition to meeting the requirements of § 1421.228 of the continuing supplement, farmers' stock peanuts to be eligible for farm-stored loan and purchase must be free of Aspergillus flavus mold as determined by a Federal-State inspector.

Effective date: September 5, 1975.

Signed at Washington, D.C., on August 27, 1975.

KENNETH E. FRICK,
Executive Vice President, Commodity Credit Corporation.

[FR Doc.75-23592 Filed 9-4-75;8:45 am]

[1] The price for all Valencia-type peanuts in the Southeast and Virginia-Carolina areas and for those Valencia-type peanuts in the Southwest area which are not suitable for cleaning and roasting will be the same as for Spanish-type peanuts in the same area.

[CCC Grain Price Support Regulations, 1975 Crop Honey Supplement]

PART 1434—HONEY

Purchase Rates

On January 15, 1975, notice of proposed rulemaking regarding purchase rates for 1975 crop honey and detailed operating provisions to carry out the 1975 crop honey purchase program was published in the FEDERAL REGISTER (40 FR 2726).

Two responses were received, one from an interested individual honey producer and the other from a marketing association. The responses recommended that a proposed purchase program permit producers to make declaration of intent to sell honey to CCC through March 31 for the previous year's production, that producers be notified of their delivery date and destination by June 1, and that delivery be made by producers on or about July 1. Specific delivery dates are not being adopted by CCC because of their impracticability under the purchase program.

The regulations contained in 7 CFR 1434.40 through 1434.44 are revised to read as follows, effective as to 1975 crop honey. The material previously appearing in these sections remains in full force and effect as to the crops to which it was applicable.

Subpart—1975 Crop Honey Purchase Program

Sec.
1434.40 Purpose.
1434.41 Availability.
1434.42 Purchase rates.
1434.43 Discounts.

AUTHORITY: Secs. 4 and 5, 62 Stat. 1070, as amended (15 U.S.C. 714 b and c); secs. 201, 401, 63 Stat. 1052, 1054 (7 U.S.C. 1446, 1421).

Subpart—1975 Crop Honey Purchase Program

§ 1434.40 Purpose.

This subpart contains program provisions which, together with (a) the Honey Purchase Regulations for 1975 and Subsequent Crops (40 FR 30798), (b) the Cooperative Marketing Association Eligibility Requirements for Price Support in Part 1425 of this chapter, and (c) any amendments to such regulations set forth the requirements with respect to purchases of 1975-crop honey.

§ 1434.41 Availability.

Producers desiring to offer eligible honey for purchase must complete a pur-

chase agreement (Form CCC-614) at the county ASCS office on or before March 31, 1976.

§ 1434.42 Purchase rates.

(a) *Table and nontable honey.* The rate for the quantity of 1975-crop honey purchased shall be the rate for the respective class and color set forth below:

Class and color: Table honey:	Cents per pound
1. White and lighter	26.3
2. Extra light amber	25.3
3. Light amber	24.3
4. Other table honey	22.3
Nontable honey	22.3

(b) *Objectionable flavor, fermentation, or caramelization.* The settlement value for a lot of honey delivered for purchase which grades substandard on account of objectionable flavor, fermentation, or caramelization shall be the lower of its market value as determined by CCC or a value determined on the basis of the purchase rate for nontable honey.

(c) *Grade not certified.* The settlement value for a lot of honey for purchase on which the grade cannot be certified shall be the lower of its market value as determined by CCC or a value as determined on the basis of the purchase rate for nontable honey.

(d) *Substandard.* The rate for a lot of honey delivered for purchase which grades substandard on account of defects or moisture or a combination of defects and moisture shall be adjusted by the discounts in § 1434.43.

§ 1434.43 Discounts.

(a) *Defects.* The purchase rate for a lot of honey delivered for purchase which grades substandard on account of defects shall be adjusted by the following discount:

Substandard account of:	Discount (cents per pound)
Defects	2

(b) *Moisture.* The purchase rate for a lot of honey delivered for purchase which contains moisture in excess of 18.5 percent shall be adjusted by the following discounts which shall be in addition to the discount for defects:

Moisture (percent):	Discount (cents per pound)
18.5	$0.0
19.0	.5
19.5	1.0
20.0	1.5
20.5	2.0
21.0	2.5
21.5	3.0
22.0	3.5
22.5	4.0
23.0	4.5
23.5	5.0
24.0	5.5
24.5	6.0

(c) *Commingled storage.* The purchase rate for a lot of honey tendered for purchase by CCC while stored commingled in a warehouse, or delivered to a warehouse in bulk, shall be adjusted by the following discount:

	Discount (cents per pound)
Bulk commingled	1.5

Effective date: September 5, 1975.

Signed at Washington, D.C., on August 27, 1975.

KENNETH E. FRICK,
Executive Vice President,
Commodity Credit Corporation.

[FR Doc.75-23593 Filed 9-4-75;8:45 am]

Title 9—Animals and Animal Products

CHAPTER I—ANIMAL AND PLANT HEALTH INSPECTION SERVICE, DEPARTMENT OF AGRICULTURE

SUBCHAPTER E—VIRUSES, SERUMS, TOXINS, AND ANALOGOUS PRODUCTS: ORGANISMS AND VECTORS

PART 113—STANDARD REQUIREMENTS

Miscellaneous Amendments

• Purpose: To make minor editorial changes for scientific accuracy in the choice of words and to correct spelling and printing errors. •

Pursuant to the authority contained in the Virus-Serum-Toxin Act of March 4, 1913 (21 U.S.C. 151-158), Part 113, Subchapter E, Chapter 1 of Title 9 of the Code of Federal Regulations is amended by making corrections as follows:

"Perfringens" was incorrectly capitalized in § 113.96 and § 113.97 and the plural of "gram" was incorrectly used in § 113.97. The word "samples" was inadvertently omitted from § 113.162(d)(3) and "per chicken" from § 113.164(e)(2)(iii). An incorrect reference was made in § 113.164(e)(3)(i). The word "centrifuge" was misprinted in § 113.202(a).

For scientific accuracy, "Potency Test" has been changed to "Virus titer requirements" and "0.7 logs" has been changed to $10^{0.7}$" where they occur in paragraphs 113.139(d)(2), 113.140(d)(2), 113.141(d)(2), 113.142(d)(3), 113.143(c)(3)(v), 113.144(d)(3), 113.145(d)(3), 113.146(d)(3), 113.160(d)(3), 113.161(d)(2), 113.162(d)(3)(iii), 113.163(d)(3), and 113.164(d)(3). Other minor editorial changes have been made for correctness, consistency, and clarity.

Also for consistency, the parenthesis has been deleted from the headings for sections 113.125, 113.128, and 113.129.

All words in the headings should be capitalized.

1. Section 113.96 is amended by revising the introductory paragraph to read:

§ 113.96 Clostridium Perfringens Type C Toxoid and Bacterin-Toxoid.

Clostridium Perfringens Type C Toxoid and Clostridium Perfringens Type C Bacterin-Toxoid shall be produced from a culture of *Clostridium perfringens* Type C which has been inactivated and is nontoxic. Each serial shall meet the applicable requirements in § 113.85 and shall be tested for purity, safety, and potency as prescribed in this section. Any serial found unsatisfactory by a prescribed test shall not be released.

* * * * *

2. Section 113.97 is amended by revising the introductory paragraph and paragraph (c)(1)(vi) to read:

§ 113.97 Clostridium Perfringens Type D Toxoid and Bacterin-Toxoid.

Clostridium Perfringens Type D Toxoid and Clostridium Perfringens Type D Bacterin-Toxoid shall be produced from a culture of *Clostridium perfringens* Type D which has been inactivated and is nontoxic. Each serial shall meet the applicable requirements in § 113.85 and shall be tested for purity, safety, and potency as prescribed in this section. Any serial found unsatisfactory by a prescribed test shall not be released.

* * * * *

(c) * * *

(1) * * *

(vi) *Diluent.* The solution used to make proper dilutions prescribed in this test. Such solutions shall be made by dissolving 1 gram of peptone and 0.25 gram of sodium chloride in each 100 ml of distilled water; adjusting the pH to 7.2; autoclaving at 250° F for 25 minutes; and storing at 4° C until used.

3. Section 113.125 is amended by revising the heading to read:

§ 113.125 Newcastle Disease Vaccine, Killed Virus.

* * * * *

§ 113.128 Avian Encephalomyelitis Vaccine, Killed Virus.

4. Section 113.128 is amended by revising the heading as set forth above.

§ 113.129 Rabies Vaccine, Killed Virus.

5. Section 113.129 is amended by revising the heading as set forth above.

6. Section 113.139 is amended by revising paragraph (d)(2) to read:

§ 113.139 Feline Panleukopenia Vaccine.

* * * * *

(d) * * *

(2) *Virus titer requirements.* Final container samples of completed product shall be tested for virus titer using the titration method used in paragraph (c)(2) of this section. To be eligible for release, each serial and each subserial shall have a virus titer sufficiently greater than the titer of vaccine virus used in the immunogenicity test prescribed in paragraph (c) of this section to assure that when tested at any time within the expiration period, each serial and subserial shall have a virus titer of $10^{0.7}$ greater than that used in such immunogenicity test but not less than $10^{3.4}$ ID_{50} per dose.

7. Section 113.140 is amended by revising paragraph (d)(2) to read:

§ 113.140 Canine Hepatitis Vaccine.

* * * * *

(d) * * *

(2) *Virus titer requirements.* Final container samples of completed product shall be tested for virus titer using the titration method used in paragraph (c)(2) of this section. To be eligible for release, each serial and each subserial shall

have a virus titer sufficiently greater than the titer of vaccine virus used in the immunogenicity test prescribed in paragraph (c) of this section to assure that when tested at any time within the expiration period, each serial and subserial shall have a virus titer of $10^{0.7}$ greater than that used in such immunogenicity test but not less than $10^{2.5}$ $TCID_{50}$ per dose.

8. Section 113.141 is amended by revising paragraph (d)(2) to read:

§ 113.141 Canine Distemper Vaccine, Ferret Avirulent.

* * * * *

(d) * * *

(2) *Virus titer requirements.* Final container samples of completed product shall be tested for virus titer using the titration method used in paragraph (c)(2) of this section. To be eligible for release, each serial and each subserial shall have a virus titer sufficiently greater than the titer of vaccine virus used in the immunogenicity test prescribed in paragraph (c) of this section to assure that when tested at any time within the expiration period, each serial and subserial shall have a virus titer of $10^{0.7}$ greater than that used in such immunogenicity test but not less than $10^{2.5}$ ID_{50} per dose.

9. Section 113.142 is amended as follows:

§ 113.142 Canine Distemper Vaccine, Ferret Virulent.

* * * * *

(d) * * *

(3) *Virus titer requirements.* Final container samples of completed product shall be tested for virus titer. To be eligible for release, each serial and subserial shall have a virus titer sufficiently greater than the virus dose used in the immunigenicity test prescribed in paragraph (c) of this section to assure that when tested at any time within the expiration period, each serial and subserial shall have a virus titer of $10^{0.7}$ greater than that used in such immunogenicity test when tested by the ferret injection method.

10. Section 113.143 is amended by revising paragraph (c)(3)(v) to read:

§ 113.143 Encephalomyelitis Vaccine, Venezuelan.

* * * * *

(c) * * *

(3) * * *

(v) Final container samples of completed product shall be tested for virus titer. To be eligible for release, each serial of vaccine shall have a $GPIPID_{50}$ titer of sufficiently greater than the titer of vaccine virus used in the immunogenicity test prescribed in paragraph (b) of this section to assure that when tested at any time within the expiration period, each serial shall have a $GPIPID_{50}$ titer $10^{0.7}$ greater than that used in such immunogenicity test but not less than $10^{3.7}$ $GPIPID_{50}$ per dose.

11. Section 113.144 is amended by revising paragraph (d)(3) to read:

§ 113.144 Bovine Parainfluenza$_3$ Vaccine.

* * * * *

(d) * * *

(3) *Virus titer requirements.* Final container samples of completed product shall be tested for virus titer using the titration method used in paragraph (c)(2) of this section. To be eligible for release, each serial and each subserial shall have a virus titer sufficiently greater than the titer of vaccine virus used in the immunogenicity test prescribed in paragraph (c) of this section to assure that when tested at any time within the expiration period, each serial and subserial shall have a virus titer of $10^{0.7}$ greater than that used in such immunogenicity test but not less than $10^{2.7}$ $TCID_{50}$ per dose.

12. Section 113.145 is amended by revising paragraph (d)(3) to read:

§ 113.145 Bovine Rhinotracheitis Vaccine.

* * * * *

(d) * * *

(3) *Virus titer requirements.* Final container samples of completed product shall be tested for virus titer using the titration method used in paragraph (c)(2) of this section. To be eligible for release, each serial and each subserial shall have a virus titer sufficiently greater than the titer of vaccine virus used in the immunogenicity test prescribed in paragraph (c) of this section to assure that when tested at any time within the expiration period, each serial and subserial shall have a virus titer of $10^{0.7}$ greater than that used in such immunogenicity test but not less than $10^{3.7}$ $TCID_{50}$ per dose.

13. Section 113.146 is amended by revising paragraph (d)(3) to read:

§ 113.146 Bovine Virus Diarrhea Vaccine.

* * * * *

(d) * * *

(3) *Virus titer requirements.* Final container samples of completed product shall be tested for virus titer using the titration method used in paragraph (c)(2) of this section. To be eligible for release, each serial and each subserial shall have a virus titer sufficiently greater than the titer of vaccine virus used in the immunogenicity test prescribed in paragraph (c) of this section to assure that when tested at any time within the expiration period, each serial and subserial shall have a virus titer of $10^{0.7}$ greater than that used in such immunogenicity test but not less than $10^{3.5}$ $TCID_{50}$ per dose.

14. Section 113.160 is amended by revising paragraph (d)(3) and the introductory paragraph in (e) to read:

§ 113.160 Avian Encephalomyelitis Vaccine.

* * * * *

(d) * * *

(3) *Virus titer requirements.* Final container samples of completed product shall be tested for virus titer using the titration method used in paragraph (c)(2) of this section. To be eligible for release, each serial and each subserial shall have a virus titer sufficiently greater than the titer of vaccine virus used in the immunogenicity test prescribed in paragraph (c) of this section to assure that when tested at any time within the expiration period, each serial and subserial shall have a virus titer of $10^{0.7}$ greater than that used in such immunogenicity test but not less than $10^{2.5}$ EID_{50} per dose.

(e) Until a lot of Master Seed Virus is established as prescribed in paragraphs (a), (b), and (c) of this section, each serial and subserial shall meet the applicable requirements prescribed in § 113.135, except paragraph (c) in paragraph (d)(1) of this section, and in this paragraph.

* * * * *

15. Section 113.161 is amended by revising paragraph (d)(2) to read:

§ 113.161 Avian Pox Vaccine.

* * * * *

(d) * * *

(2) *Virus titer requirements.* Final container samples of completed product shall be tested for virus titer using the titration method used in paragraph (c)(2) of this section. To be eligible for release, each serial and each subserial shall have a virus titer sufficiently greater than the titer of vaccine virus used in the immunogenicity test prescribed in paragraph (c) of this section to assure that when tested at any time within the expiration period, each serial and subserial shall have a virus titer of $10^{0.7}$ greater than that used in such immunogenicity test but not less than $10^{2.0}$ EID_{50} per dose.

16. Section 113.162 is amended by revising the introductory portion of paragraph (d)(3) and paragraph (d)(3)(iii) to read:

§ 113.162 Bronchitis Vaccine.

* * * * *

(d) * * *

(3) *Virus titer requirements.* Final container samples of completed product shall be tested for virus titer using the procedure prescribed in paragraph (c)(2) of this section and in this paragraph.

* * * * *

(iii) To be eligible for release, each serial and each subserial shall have a virus titer sufficiently greater than the titer of vaccine virus used in the immunogenicity test prescribed in paragraph (c) of this section to assure that when tested at any time within the expiration period, each serial and subserial shall have a virus titer of $10^{0.7}$ greater than that used in such immunogenicity test but not less than $10^{2.0}$ EID_{50} per dose.

17. Section 113.163 is amended by revising paragraph (d)(3) to read:

§ 113.163 Fowl Laryngotracheitis Vaccine.

* * * * *

(d) * * *

(3) *Virus titer requirements.* Final container samples of completed product

shall be tested for virus titer using the titration method provided in paragraphs (c) (2) or (3) of this section. To be eligible for release, each serial and each subserial shall have a virus titer sufficiently greater than the titer of vaccine virus used in the immunogenicity test prescribed in paragraph (c) of this section to assure that when tested at any time within the expiration period, each serial and subserial shall have a virus titer of $10^{0.7}$ greater than that used in such immunogenicity test but not less than $10^{2.5}$ EID_{50} per dose for chicken embryo origin vaccine and $10^{2.5}$ EID_{50} or $10^{2.5}$ $TCID_{50}$ per dose for tissue culture origin vaccine.

* * * * *

18. Section 113.164 is amended by revising paragraphs (b), (d)(3), (e)(1), (e)(2)(iii), and (e)(3)(i) to read:

§ 113.164 Newcastle Disease Vaccine.

* * * * *

(b) Each lot of Master Seed Virus shall be tested for pathogens by the chicken embryo inoculation test prescribed in § 113.37, except that, if the test is inconclusive because of a vaccine virus override, the test may be repeated and if the repeat test is inconclusive for the same reason, the chicken inoculation test prescribed in § 113.36 may be conducted and the virus judged accordingly.

* * * * *

(d) * * *

(3) *Virus titer requirements.* Final container samples of completed product shall be tested for virus titer using the titration method used in paragraph (c)(2) of this section. To be eligible for release, each serial and each subserial shall have a virus titer sufficiently greater than the titer of vaccine virus used in the immunogenicity test prescribed in paragraph (c) of this section to assure that when tested at any time within the expiration period, each serial and subserial shall have a virus titer of $10^{0.7}$ greater than that used in such immunogenicity test but not less than $10^{2.5}$ EID_{50} per dose.

(e) * * *

(1) A virus titration shall be conducted on final container samples of completed product in accordance with the titration methods used in paragraph (c)(2) of this section. A serial or subserial which does not contain at least $10^{2.5}$ EID_{50} per dose of Newcastle disease virus through the expiration date is unsatisfactory.

(2) * * *

(iii) Twenty to twenty-eight days post-vaccination, the vaccinates and the controls shall be challenged intramuscularly with at least $10^{4.5}$ EID_{50} per chicken of Newcastle disease virus provided or approved by Veterinary Services. The chickens shall be observed each day for 14 days.

* * * * *

(3) * * *

(i) Vaccines recommended for use in chickens 10 days of age or younger shall be tested in accordance with paragraphs (d)(2)(i), (ii), and (iii) of this section.

* * * * *

(37 Stat. 832-833; 21 U.S.C. 151-158)

These amendments are administrative or editorial and the changes are corrective or conformative in nature and make no substantive changes in the affected regulations.

Accordingly, under the administrative procedure provisions in 5 U.S.C. 553, it is found upon good cause that notice and other public procedure concerning the amendments are impracticable and unnecessary, and good cause is found for making the amendments effective less than 30 days after publication in the FEDERAL REGISTER.

The foregoing amendments shall become effective upon issuance.

Done at Washington, DC, this 29th day of August 1975.

PIERRE A. CHALOUX,
Acting Deputy Administrator, Veterinary Services, Animal and Plant Health Inspection Service.

[FR Doc.75-23591 Filed 9-4-75;8:45 am]

Title 12—Banks and Banking

CHAPTER VII—NATIONAL CREDIT UNION ADMINISTRATION

PART 701—ORGANIZATION AND OPERATION OF FEDERAL CREDIT UNIONS

Cashing Checks and Money Orders

On pages 30291-30292 of the July 18, 1975, edition of the FEDERAL REGISTER (40 FR 30291-30292) there was published a proposal to amend Part 701 (12 CFR 701) by revising § 701.23(d). The purpose of the proposal is to give to each Federal credit union the option of charging a fee when the cashing of a check or money order is not applied in its entirety for payment of a loan, payment of interest, payment of any obligation to the credit union, or the purchase of shares. Interested persons were given until August 11, 1975, to submit written comments, suggestions and objections regarding the proposed amendment. As a result of the comments, no changes were deemed necessary.

Accordingly, the proposed amendment to Part 701 (12 CFR 701) is adopted as set forth below and is effective immediately.

HERMAN NICKERSON, Jr.,
Administrator.

AUGUST 28, 1975.

(Sec. 120, 73 Stat. 635, (12 U.S.C. 1766) and Sec. 209, 84 Stat. 1014, (12 U.S.C. 1789))

§ 701.23 Cashing checks and money orders.

* * * * *

(d) No fee shall be charged by a Federal credit union to a member for the cashing of a check or money order when such check or money order is applied in its entirety for payment of a loan, payment of interest, payment of any obligation to the credit union, or the purchase of shares. Nor shall any fee be charged to the member for the cashing of a check or money order drawn by the Federal credit union on its own bank account and issued to the member in connection with a withdrawal by the member from a share account or in connection with the disbursement of a loan.

[FR Doc.75-23562 Filed 9-4-75;8:45 am]

Title 14—Aeronautics and Space

CHAPTER I—FEDERAL AVIATION ADMINISTRATION, DEPARTMENT OF TRANSPORTATION

[Airworthiness Docket No. 75-WE-1-AD; Amdt. 39-2357]

PART 39—AIRWORTHINESS DIRECTIVES

AiResearch Model GTCP660-4 and -4R Auxiliary Power Units (APU)

Pursuant to a Notice of Proposed Rule Making, published June 12, 1975, in the FEDERAL REGISTER, (40 FR 25027), the agency proposed to amend Amendment 39-2108 (40 FR 8541), AD 75-05-17, as amended by Amendment 39-2224 (40 FR 23722), to require accomplishment of modifications and inspections described in AiResearch Service Bulletin GTCP 660-49-3713, dated May 15, 1975, or later FAA-approved revisions, within six months of the effective date of the adopted rule on certain Boeing 747 aircraft. When the modifications and inspections were performed, the placard required by paragraph (a) of the AD may be removed, and the inspections required by paragraph (b) may be discontinued.

The agency considered the incorporation of these modifications and inspections as necessary to minimize the hazard of injury to persons near the aircraft during ground operations.

Interested persons have been afforded an opportunity to participate in the making of the amendment.

Three comments were received. The Air Transport Association on behalf of member air carriers, British Airways, and Belgian World Airlines (Sabena), as Central Agency for the ATLAS group (Air France, Lufthansa, Alitalia, Iberia and Sabena) submitted comments. The comments stressed problems of parts availability, scheduling capabilities of the operators, and spare APU availability; Sabena also noted that there had been only one penetration of the aircraft Body Station 2658 firewall due to non-containment of fragments during over more than 2,900,000 total aircraft hours of operation. Based on the foregoing, the comments requested substantial extensions of the compliance time within which the actions were to be taken in the adopted rule, while concurring in the agency's intent to effect a terminating action with some time frame.

The Sabena comment referencing the incident in which there had been penetration of the firewall at Body Station 2658 is true, but is not pertinent to the agency's intent to protect ground personnel from shrapnel piercing the adjacent fuselage. In addition to the fragments which pierced the forward firewall of the APU compartment during the incident cited above, there were numerous fragments which penetrated the lower sur-

face of the APU compartment and could have endangered ground personnel and produced a condition which could impair fire containment because of firewall degradation.

With respect to the compliance time proposed in the Notice of Proposed Rule Making, it is to be noted that the agency previously, by Amendment 39-2224, referenced above, issued on May 21, 1975, and made effective on June 5, 1975, authorized the operators to incorporate the modifications and inspections of the manufacturer's service bulletin as a terminating action to the placard requirement of paragraph (a) of the AD, and the inspections of paragraph (b). Amendment 39-2224 was adopted as an immediately adopted rule, as it provided an alternative means of compliance, which, when performed, relieved the operators of the AD requirements, as indicated above. The agency believes that the operators, both in the interest of safety for the ground personnel involved, and the advantages of eliminating the restrictions of the AD, would have been incorporating the actions set forth in the service bulletin on a scheduled basis even before the issuance of the Notice. And, while it is true that there has been but one incident involving the failure to contain the fragments of the APU, the potential hazard presented by the occurrence is sufficiently serious to warrant effective corrective action at the earliest time, and on a practicable basis.

The ATA comment suggested a compliance time of 24 months or 1,000 hours time in service on the APU, whichever occurs later.

British Airways suggested a date of June 30, 1976. Sabena suggested that the service bulletin be incorporated the first time the APU is removed from the airplane with a maximum delay of 24 months after the effective date of the amendment.

The agency has considered these comments. In the interest of safety in aviation, the agency believes that the compliance time adopted herein, namely, on or before July 1, 1976, to do the work, represents a reasonable standard.

In consideration of the foregoing, and pursuant to the authority delegated to me by the Administrator (31 FR 13697), § 39.13 of Part 39 of the Federal Aviation Regulations, Amendment 39-2108 (40 FR 8541), AD 75-05-17, as amended by Amendment 39-2224 (40 FR 23722), is further amended, by revising paragraph (c) to read:

§ 39.13 [Amended]

(c) The modifications and inspections described in AiResearch Bulletin GTCP 660-49-3713, dated May 15, 1975, or later FAA-approved revisions must be incorporated on or before July 1, 1976. Upon completion of these modifications and inspections, the placard required by paragraph (a), above, may be removed and the inspections required by (b), above, may be discontinued.

This amendment becomes effective October 6, 1975.

This amendment is made under the authority of sections 313(a), 601, and 603 of the Federal Aviation Act of 1958 (49 U.S.C. 1354(a), 1421, and 1423) and of section 6(c) of the Department of Transportation Act (49 U.S.C. 1655(c)).

Issued in Los Angeles, California on August 21, 1975.

LYNN L. HINK,
Acting Director,
FAA Western Region.

[FR Doc.75-23488 Filed 9-4-75;8:45 am]

[Airworthiness Docket No. 75-SW-11, Amdt. 39-2350]

PART 39—AIRWORTHINESS DIRECTIVES

Bell Models 206A, 206B, 206A-1, and 206B-1 Helicopters

Amendment 39-2122 (40 FR 10661), AD 75-06-03, as revised by Amendment 39-2146 (40 FR 14297) requires an immediate and a 100-hour repetitive inspection for possible cracks in the upper and lower clevis on each main rotor blade pitch link assembly, P/N 206-010-330 or 206-010-342, and an inspection of the outer swashplate ring horn bearings for excessive breakaway torque on Bell Models 206A, 206B, 206A-1 and 206B-1 helicopters. After issuing Amendment 39-2122, AD 75-06-03, and Amendment 39-2146, the agency has been informed of satisfactory service history on the affected clevises and helicopters by several operators and by the Helicopter Association of America. Bell Helicopter Company has inspected approximately 40 clevises with alleged cracks that were returned to them as prescribed in the Mailgram dated February 15, 1975, from Bell Helicopter Company to all 206A, 206B, and TH57A operators. None of these clevises had cracks in the threaded shanks. Amendment 39-2122, AD 75-06-03, has been in effect since March 12, 1975, and the FAA has not received a confirmed case of a cracked clevis as a result of this AD. Based on this information and the numerous requests for relief from further repetitive inspections, the AD is being amended to delete any further requirement for a repetitive inspection.

The agency again requests interested persons to submit written information and comments on the main rotor pitch link assemblies as they may desire. Communications should identify the docket number and be submitted in triplicate to the Regional Counsel, Southwest Region, Federal Aviation Administration, P.O. Box 1689, Forth Worth, Texas 76101. All comments will be available in the Office of Regional Counsel for examination by interested persons.

The comments received on or before September 22, 1975, that are submitted in response to the request for additional information will be considered by the Director and the amendment may be changed in the light of those comments prior to the adoption on October 2, 1975. Since this amendment provides relief from further repetitive inspections and imposes no additional burden on any person, the amendment may be made effective in 40 days.

In consideration of the foregoing, and pursuant to the authority delegated to me by the Administrator (31 FR 13697), Section 39.13 of Part 39 of the Federal Aviation Regulations, Amendment 39-2122 (40 FR 10661), AD 75-06-03 as amended by Amendment 39-2146 (40 FR 14297) is further amended by changing the compliance paragraph to read as follows:

Compliance required within 10 hours' time in service after March 12, 1975, unless already accomplished.

This amendment becomes effective October 2, 1975.

This amendment is made under the authority of Sections 313(a), 601, and 603 of the Federal Aviation Act of 1958 (49 U.S.C. 1354(a), 1421, and 1423) and of Section 6(c) of the Department of Transportation Act (49 U.S.C. 1655(c)).

Issued in Fort Worth, Texas, on August 13, 1975.

HENRY L. NEWMAN,
Director, Southwest Region.

[FR Doc.75-23489 Filed 9-4-75;8:45 am]

[Airworthiness Docket No. 75-SW-39; Amdt. 39-2356]

PART 39—AIRWORTHINESS DIRECTIVES

Rockwell Models 690, 690A, and 685

Amendment 39-2275, 40 FR 29272, AD 75-15-01, imposes a speed and deflection limitation during flap operation, and requires a one-time inspection of flap bracket, P/N 510003-92, with a subsequent modification prior to lifting the flap speed and deflection limitation. After issuing Amendment 39-2257, service experience established that cracks were still occurring while operating with the flap speed and deflection limitation.

Therefore, the AD is being superseded by a new AD that requires a repetitive inspection while operating in accordance with the flap speed and deflection limitation. Also, further modification information is provided.

Since a situation exists that requires immediate adoption of this regulation, it is found that notice and public procedure hereon are impracticable and good cause exists for making this amendment effective in less than 30 days.

In consideration of the foregoing, and pursuant to the authority delegated to me by the Administrator (31 FR 13697), § 39.13 of Part 39 of the Federal Aviation Regulations is amended by adding the following new airworthiness directive:

ROCKWELL: Applies to Models 690, S/N 11000 and subsequent, 690A, S/N 11100 and subsequent, and 685, S/N 12000 and subsequent.

Compliance required as indicated:

A. Immediately upon receipt of this AD, limit flap speed and deflection as follows: