from the production and marketing situation confronting the lemon industry.

(i) The committee has submitted its recommendation with respect to the quantity of lemons it deems advisable to be handled during the ensuing week. Such recommendation resulted from consideration of the factors enumerated in the order. The committee further reports the demand for lemons is good on all sizes and grades, except for size 165's. Average f.o.b. price was $4.99 per carton the week ended December 28, 1974, compared to $4.72 per carton the previous week. Track and rolling supplies at 102 cars were down 8 cars from last week.

(ii) Having considered the recommendation and information submitted by the committee, and other available information, the Secretary finds that the quantity of lemons which may be handled should be fixed as hereinafter set forth.

(3) It is hereby further found that it is impracticable and contrary to the public interest to give preliminary notice, engage in public rule-making procedure, and postpone the effective date of this section until 30 days after publication hereof in the FEDERAL REGISTER (5 U.S.C. 553) because the time intervening between the date when information upon which this section is based became available and the time when this section must become effective in order to effectuate the declared policy of the act is insufficient, and a reasonable time is permitted, under the circumstances, for preparation for such effective time; and good cause exists for making the provisions hereof effective as hereinafter set forth. The committee held an open meeting during the current week, after giving due notice thereof, to consider supply and market conditions for lemons and the need for regulation; interested persons were afforded an opportunity to submit information and views at this meeting; the recommendation and supporting information for regulation during the period specified herein were promptly submitted to the Department after such meeting was held; the provisions of this section, including its effective time, are identical with the aforesaid recommendation of the committee, and information concerning such provisions and effective time has been disseminated among handlers of such lemons; it is necessary, in order to effectuate the declared policy of the act, to make this section effective during the period herein specified; and compliance with this section will not require any special preparation on the part of persons subject hereto which cannot be completed on or before the effective date hereof. Such committee meeting was held on December 30, 1974.

(b) *Order.* (1) The quantity of lemons grown in California and Arizona which may be handled during the period January 5, 1975, through January 11, 1975, is hereby fixed at 195,000 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: December 31, 1974.

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

[FR Doc.75-219 Filed 1-2-75;8:45 am]

Title 31—Fiscal Service

SUBCHAPTER B—BUREAU OF THE PUBLIC DEBT

PART 316—OFFERING OF UNITED STATES SAVINGS BONDS

Series E

The purpose of this first supplement to Department of the Treasury Circular No. 653, Ninth Revision, dated March 18, 1974 (31 CFR Part 316), is to show the redemption values and investment yields for the next extended maturity period for United States Savings Bonds of Series E bearing issue dates of (1) June 1 through November 1, 1945, (2) June 1 through September 1, 1955, (3) October 1 through November 1, 1955, (4) June 1 through November 1, 1968, and (5) June 1 through November 1, 1969. Accordingly, in § 316.14 the tables to the circular are hereby supplemented by the addition of Tables 12-A, 39-A, 40-A, 86-A and 88-A.

Dated: December 24, 1974.

JOHN K. CARLOCK,
Fiscal Assistant Secretary.

§ 316.14 Reservations as to terms of offer.

* * * * *

TABLE 12

BONDS BEARING ISSUE DATES FROM JUNE 1 THROUGH NOV. 1, 1945

Issue price	$7.50	$18.75	$37.50	$75.00	$150.00	$375.00	$750.00	Approximate investment yield (annual percentage rate)		
Denomination	10.00	25.00	50.00	100.00	200.00	500.00	1000.00			
Period (years and months after second extended maturity at 30 years 0 months)	(1) Redemption values during each half-year period (values increase on first day of period)*							(2) From beginning of current maturity period to beginning of each ½-yr. pd.	(3) From beginning of each ½-yr. period to beginning of next ½-yr. pd.	(4) From beginning of each ½-yr. period to 3rd extended maturity
	THIRD EXTENDED MATURITY PERIOD**									
								Percent	Percent	Percent
0-0 to 0-6	$22.46	$56.15	$112.30	$224.60	$449.20	$1123.00	$2246.00	----	5.98	6.00
0-6 to 1-0	23.13	57.83	115.66	231.32	462.64	1156.60	2313.20	5.98	6.02	6.00
1-0 to 1-6	23.83	59.57	119.14	238.28	476.56	1191.40	2382.80	6.00	6.01	6.00
1-6 to 2-0	24.54	61.36	122.72	245.44	490.88	1227.20	2454.40	6.00	6.00	6.00
2-0 to 2-6	25.28	63.20	126.40	252.80	505.60	1264.00	2528.00	6.00	5.98	6.00
2-6 to 3-0	26.04	65.09	130.18	260.36	520.72	1301.80	2603.60	6.00	6.02	6.00
3-0 to 3-6	26.82	67.05	134.10	268.20	536.40	1341.00	2682.00	6.00	6.00	6.00
3-6 to 4-0	27.62	69.06	138.12	276.24	552.48	1381.20	2762.40	6.00	5.99	6.00
4-0 to 4-6	28.45	71.13	142.26	284.52	569.04	1422.60	2845.20	6.00	5.99	6.00
4-6 to 5-0	29.30	73.26	146.52	293.04	586.08	1465.20	2930.40	6.00	6.01	6.00
5-0 to 5-6	30.18	75.46	150.92	301.84	603.68	1509.20	3018.40	6.00	5.99	6.00
5-6 to 6-0	31.09	77.72	155.44	310.88	621.76	1554.40	3108.80	6.00	6.02	6.00
6-0 to 6-6	32.02	80.06	160.12	320.24	640.48	1601.20	3202.40	6.00	6.00	6.00
6-6 to 7-0	32.98	82.46	164.92	329.84	659.68	1649.20	3298.40	6.00	5.99	6.00
7-0 to 7-6	33.97	84.93	169.86	339.72	679.44	1698.60	3397.20	6.00	6.00	6.00
7-6 to 8-0	34.99	87.48	174.96	349.92	699.84	1749.60	3499.20	6.00	5.99	6.00
8-0 to 8-6	36.04	90.10	180.20	360.40	720.80	1802.00	3604.00	6.00	6.02	6.00
8-6 to 9-0	37.12	92.81	185.62	371.24	742.48	1856.20	3712.40	6.00	5.99	6.00
9-0 to 9-6	38.24	95.59	191.18	382.36	764.72	1911.80	3823.60	6.00	6.00	6.00
9-6 to10-0	39.38	98.46	196.92	393.84	787.68	1969.20	3938.40	6.00	5.99	5.99
10-0 2/	40.56	101.41	202.82	405.64	811.28	2028.20	4056.40	6.00 3/	----	----

1/ Month, day, and year on which issues of June 1, 1945, enter each period. For subsequent issue months add the appropriate number of months.
2/ Third extended maturity value reached at 40 years and 0 months after issue.
3/ Yield on purchase price from issue date to third extended maturity date is 4.26 percent.

* For earlier redemption values and yields see appropriate table in Department Circular 653, 9th Revision, as amended and supplemented.
** This table does not apply if the prevailing rate for Series K bonds being issued at the time the extension begins is different from 6.00 percent.

TABLE 39

BONDS BEARING ISSUE DATES FROM JUNE 1 THROUGH SEPT. 1, 1955

Issue price	$18.75	$37.50	$75.00	$150.00	$375.00	$750.00	$7500	Approximate investment yield (annual percentage rate)
Denomination	25.00	50.00	100.00	200.00	500.00	1000.00	10000	

Period (years and months after first extended maturity at 19 years 8 months)	(1) Redemption values during each half-year period (values increase on first day of period)* SECOND EXTENDED MATURITY PERIOD**							(2) From beginning of current maturity period to beginning of each ½-yr. pd.	(3) From beginning of each ½-yr. period to beginning of next ½-yr. pd.	(4) From beginning of each ½-yr. period to 2nd extended maturity
								Percent	Percent	Percent
0-0 to 0-6 . . . 1/ (2/1/75)	$40.93	$ 81.86	$163.72	$327.44	$ 818.60	$1637.20	$16372	----	6.01	6.00
0-6 to 1-0 . . . . (8/1/75)	42.16	84.32	168.64	337.28	843.20	1686.40	16864	6.01	5.98	6.00
1-0 to 1-6 . . . . (2/1/76)	43.42	86.84	173.68	347.36	868.40	1736.80	17368	5.99	6.03	6.00
1-6 to 2-0 . . . . (8/1/76)	44.73	89.46	178.92	357.84	894.60	1789.20	17892	6.01	5.99	6.00
2-0 to 2-6 . . . . (2/1/77)	46.07	92.14	184.28	368.56	921.40	1842.80	18428	6.00	5.99	6.00
2-6 to 3-0 . . . . (8/1/77)	47.45	94.90	189.80	379.60	949.00	1898.00	18980	6.00	5.99	6.00
3-0 to 3-6 . . . . (2/1/78)	48.87	97.74	195.48	390.96	977.40	1954.80	19548	6.00	6.02	6.00
3-6 to 4-0 . . . . (8/1/78)	50.34	100.68	201.36	402.72	1006.80	2013.60	20136	6.00	6.00	6.00
4-0 to 4-6 . . . . (2/1/79)	51.85	103.70	207.40	414.80	1037.00	2074.00	20740	6.00	5.98	6.00
4-6 to 5-0 . . . . (8/1/79)	53.40	106.80	213.60	427.20	1068.00	2136.00	21360	6.00	6.03	6.00
5-0 to 5-6 . . . . (2/1/80)	55.01	110.02	220.04	440.08	1100.20	2200.40	22004	6.00	6.00	6.00
5-6 to 6-0 . . . . (8/1/80)	56.66	113.32	226.64	453.28	1133.20	2266.40	22664	6.00	6.00	6.00
6-0 to 6-6 . . . . (2/1/81)	58.36	116.72	233.44	466.88	1167.20	2334.40	23344	6.00	6.00	6.00
6-6 to 7-0 . . . . (8/1/81)	60.11	120.22	240.44	480.88	1202.20	2404.40	24044	6.00	5.99	6.00
7-0 to 7-6 . . . . (2/1/82)	61.91	123.82	247.64	495.28	1238.20	2476.40	24764	6.00	6.01	6.00
7-6 to 8-0 . . . . (8/1/82)	63.77	127.54	255.08	510.16	1275.40	2550.80	25508	6.00	5.99	6.00
8-0 to 8-6 . . . . (2/1/83)	65.68	131.36	262.72	525.44	1313.60	2627.20	26272	6.00	6.00	6.00
8-6 to 9-0 . . . . (8/1/83)	67.65	135.30	270.60	541.20	1353.00	2706.00	27060	6.00	6.00	6.00
9-0 to 9-6 . . . . (2/1/84)	69.68	139.36	278.72	557.44	1393.60	2787.20	27872	6.00	6.00	6.00
9-6 to10-0 . . . . (8/1/84)	71.77	143.54	287.08	574.16	1435.40	2870.80	28708	6.00	5.99	5.99
10-0 2/ . . . . . (2/1/85)	73.92	147.84	295.68	591.36	1478.40	2956.80	29568	6.00 3/	----	----

1/ Month, day, and year on which issues of June 1, 1955, enter each period. For subsequent issue months add the appropriate number of months.
2/ Second extended maturity value reached at 29 years 8 months after issue.
3/ Yield on purchase price from issue date to 2nd extended maturity date is 4.68 percent.

* For earlier redemption values and yields see appropriate table in Department Circular 653, 9th Revision, as amended and supplemented.
** This table does not apply if the prevailing rate for Series E bonds being issued at the time the extension begins is different from 6.00 percent.

TABLE 40

BONDS BEARING ISSUE DATE OCT. 1 OR NOV. 1, 1955

Issue price	$18.75	$37.50	$75.00	$150.00	$375.00	$750.00	$7500	Approximate investment yield (annual percentage rate)
Denomination	25.00	50.00	100.00	200.00	500.00	1000.00	10000	

Period (years and months after first extended maturity at 19 years 8 months)	(1) Redemption values during each half-year period (values increase on first day of period)* SECOND EXTENDED MATURITY PERIOD**							(2) From beginning of current maturity period to beginning of each ½-yr. pd.	(3) From beginning of each ½-yr. period to beginning of next ½-yr. pd.	(4) From beginning of each ½-yr. period to 2nd extended maturity
								Percent	Percent	Percent
0-0 to 0-6 . . . 1/ (6/1/75)	$41.38	$ 82.76	$165.52	$331.04	$ 827.60	$1655.20	$16552	----	5.99	6.00
0-6 to 1-0 . . . . (12/1/75)	42.62	85.24	170.48	340.96	852.40	1704.80	17048	5.99	6.01	6.00
1-0 to 1-6 . . . . (6/1/76)	43.90	87.80	175.60	351.20	878.00	1756.00	17560	6.00	6.01	6.00
1-6 to 2-0 . . . . (12/1/76)	45.22	90.44	180.88	361.76	904.40	1808.80	18088	6.00	5.97	6.00
2-0 to 2-6 . . . . (6/1/77)	46.57	93.14	186.28	372.56	931.40	1862.80	18628	6.00	6.01	6.00
2-6 to 3-0 . . . . (12/1/77)	47.97	95.94	191.88	383.76	959.40	1918.80	19188	6.00	6.00	6.00
3-0 to 3-6 . . . . (6/1/78)	49.41	98.82	197.64	395.28	988.20	1976.40	19764	6.00	5.99	6.00
3-6 to 4-0 . . . . (12/1/78)	50.89	101.78	203.56	407.12	1017.80	2035.60	20356	6.00	6.01	6.00
4-0 to 4-6 . . . . (6/1/79)	52.42	104.84	209.68	419.36	1048.40	2096.80	20968	6.00	5.99	6.00
4-6 to 5-0 . . . . (12/1/79)	53.99	107.98	215.96	431.92	1079.80	2159.60	21596	6.00	6.00	6.00
5-0 to 5-6 . . . . (6/1/80)	55.61	111.22	222.44	444.88	1112.20	2224.40	22244	6.00	6.01	6.00
5-6 to 6-0 . . . . (12/1/80)	57.28	114.56	229.12	458.24	1145.60	2291.20	22912	6.00	6.01	6.00
6-0 to 6-6 . . . . (6/1/81)	59.00	118.00	236.00	472.00	1180.00	2360.00	23600	6.00	6.00	6.00
6-6 to 7-0 . . . . (12/1/81)	60.77	121.54	243.08	486.16	1215.40	2430.80	24308	6.00	5.99	6.00
7-0 to 7-6 . . . . (6/1/82)	62.59	125.18	250.36	500.72	1251.80	2503.60	25036	6.00	6.01	6.00
7-6 to 8-0 . . . . (12/1/82)	64.47	128.94	257.88	515.76	1289.40	2578.80	25788	6.00	5.99	6.00
8-0 to 8-6 . . . . (6/1/83)	66.40	132.80	265.60	531.20	1328.00	2656.00	26560	6.00	5.99	6.00
8-6 to 9-0 . . . . (12/1/83)	68.39	136.78	273.56	547.12	1367.80	2735.60	27356	6.00	6.02	6.01
9-0 to 9-6 . . . . (6/1/84)	70.45	140.90	281.80	563.60	1409.00	2818.00	28180	6.00	5.99	6.00
9-6 to10-0 . . . . (12/1/84)	72.56	145.12	290.24	580.48	1451.20	2902.40	29024	6.00	6.01	6.01
10-0 2/ . . . . . (6/1/85)	74.74	149.48	298.96	597.92	1494.80	2989.60	29896	6.00 3/	----	----

1/ Month, day, and year on which issues of Oct. 1, 1955, enter each period. For issues of Nov. 1, 1955, add 1 month.
2/ Second extended maturity value reached at 29 years 8 months after issue.
3/ Yield on purchase price from issue date to 2nd extended maturity date is 4.72 percent.

* For earlier redemption values and yields see appropriate table in Department Circular 653, 9th Revision, as amended and supplemented.
** This table does not apply if the prevailing rate for Series E bonds being issued at the time the extension begins is different from 6.00 percent.

TABLE 86

BONDS BEARING ISSUE DATES FROM JUNE 1 THROUGH NOV. 1, 1968

Issue price / Denomination	$18.75 / 25.00	$37.50 / 50.00	$56.25 / 75.00	$75.00 / 100.00	$150.00 / 200.00	$375.00 / 500.00	$750.00 / 1000.00	$7500 / 10000	Approximate investment yield (annual percentage rate)

Period (years and months after original maturity at* 7 years 0 months)	(1) Redemption values during each half-year period (values increase on first day of period)* — EXTENDED MATURITY PERIOD**								(2) From beginning of current maturity period to beginning of each ½-yr. pd.	(3) From beginning of each ½-yr. period to beginning of next ½-yr. pd.	(4) From beginning of each ½-yr. period to extended maturity
									Percent	Percent	Percent
0-0 to 0-6 . . . 1/(6/1/75)	$26.81	$53.62	$80.43	$107.24	$214.48	$536.20	$1072.40	$10724	——	5.97	6.00
0-6 to 1-0 . . . . (12/1/75)	27.61	55.22	82.83	110.44	220.88	552.20	1104.40	11044	5.97	6.01	6.00
1-0 to 1-6 . . . . (6/1/76)	28.44	56.88	85.32	113.76	227.52	568.80	1137.60	11376	5.99	6.05	6.00
1-6 to 2-0 . . . . (12/1/76)	29.30	58.60	87.90	117.20	234.40	586.00	1172.00	11720	6.01	5.94	6.00
2-0 to 2-6 . . . . (6/1/77)	30.17	60.34	90.51	120.68	241.36	603.40	1206.80	12068	5.99	6.03	6.00
2-6 to 3-0 . . . . (12/1/77)	31.08	62.16	93.24	124.32	248.64	621.60	1243.20	12432	6.00	5.98	6.00
3-0 to 3-6 . . . . (6/1/78)	32.01	64.02	96.03	128.04	256.08	640.20	1280.40	12804	6.00	6.00	6.00
3-6 to 4-0 . . . . (12/1/78)	32.97	65.94	98.91	131.88	263.76	659.40	1318.80	13188	6.00	6.01	6.00
4-0 to 4-6 . . . . (6/1/79)	33.96	67.92	101.88	135.84	271.68	679.20	1358.40	13584	6.00	6.01	6.00
4-6 to 5-0 . . . . (12/1/79)	34.98	69.96	104.94	139.92	279.84	699.60	1399.20	13992	6.00	6.00	6.00
5-0 to 5-6 . . . . (6/1/80)	36.03	72.06	108.09	144.12	288.24	720.60	1441.20	14412	6.00	6.00	6.00
5-6 to 6-0 . . . . (12/1/80)	37.11	74.22	111.33	148.44	296.88	742.20	1484.40	14844	6.00	5.98	6.00
6-0 to 6-6 . . . . (6/1/81)	38.22	76.44	114.66	152.88	305.76	764.40	1528.80	15288	6.00	6.02	6.00
6-6 to 7-0 . . . . (12/1/81)	39.37	78.74	118.11	157.48	314.96	787.40	1574.80	15748	6.00	5.99	6.00
7-0 to 7-6 . . . . (6/1/82)	40.55	81.10	121.65	162.20	324.40	811.00	1622.00	16220	6.00	6.02	6.00
7-6 to 8-0 . . . . (12/1/82)	41.77	83.54	125.31	167.08	334.16	835.40	1670.80	16708	6.00	5.99	6.00
8-0 to 8-6 . . . . (6/1/83)	43.02	86.04	129.06	172.08	344.16	860.40	1720.80	17208	6.00	6.00	6.00
8-6 to 9-0 . . . . (12/1/83)	44.31	88.62	132.93	177.24	354.48	886.20	1772.40	17724	6.00	6.00	6.00
9-0 to 9-6 . . . . (6/1/84)	45.64	91.28	136.92	182.56	365.12	912.80	1825.60	18256	6.00	6.00	6.00
9-6 to10-0 . . . . (12/1/84)	47.01	94.02	141.03	188.04	376.08	940.20	1880.40	18804	6.00	6.00	6.00
10-0 2/ . . . . . (6/1/85)	48.42	96.84	145.26	193.68	387.36	968.40	1936.80	19368	6.00 3/	——	——

1/ Month, day, and year on which issues of June 1, 1968, enter each period. For subsequent issue months add the appropriate number of months.
2/ Extended maturity value reached at 17 years 0 months after issue.
3/ Yield on purchase price from issue date to extended maturity date is 5.66 percent.

* For earlier redemption values and yields see appropriate table in Department Circular 653, 9th Revision, as amended and supplemented.
** This table does not apply if the prevailing rate for Series K bonds being issued at the time the extension begins is different from 6.00 percent.

TABLE 88

BONDS BEARING ISSUE DATES FROM JUNE 1 THROUGH NOV. 1, 1969

Issue price / Denomination	$18.75 / 25.00	$37.50 / 50.00	$56.25 / 75.00	$75.00 / 100.00	$150.00 / 200.00	$375.00 / 500.00	$750.00 / 1000.00	$7500 / 10000	Approximate investment yield (annual percentage rate)

Period (years and months after original maturity at 5 years 10 months)	(1) Redemption values during each half-year period (values increase on first day of period)* — EXTENDED MATURITY PERIOD**								(2) From beginning of current maturity period to beginning of each ½-yr. pd.	(3) From beginning of each ½-yr. period to beginning of next ½-yr. pd.	(4) From beginning of each ½-yr. period to extended maturity
									Percent	Percent	Percent
0-0 to 0-6 . . . 1/(4/1/75)	$25.77	$51.54	$77.31	$103.08	$206.16	$515.40	$1030.80	$10308	——	5.98	6.00
0-6 to 1-0 . . . . (10/1/75)	26.54	53.08	79.62	106.16	212.32	530.80	1061.60	10616	5.98	6.03	6.00
1-0 to 1-6 . . . . (4/1/76)	27.34	54.68	82.02	109.36	218.72	546.80	1093.60	10936	6.00	6.00	6.00
1-6 to 2-0 . . . . .(10/1/76)	28.16	56.32	84.48	112.64	225.28	563.20	1126.40	11264	6.00	5.97	6.00
2-0 to 2-6 . . . . (4/1/77)	29.00	58.00	87.00	116.00	232.00	580.00	1160.00	11600	5.99	6.00	6.00
2-6 to 3-0 . . . . (10/1/77)	29.87	59.74	89.61	119.48	238.96	597.40	1194.80	11948	5.99	6.03	6.00
3-0 to 3-6 . . . . (4/1/78)	30.77	61.54	92.31	123.08	246.16	615.40	1230.80	12308	6.00	5.98	6.00
3-6 to 4-0 . . . . (10/1/78)	31.69	63.38	95.07	126.76	253.52	633.80	1267.60	12676	6.00	6.00	6.00
4-0 to 4-6 . . . . (4/1/79)	32.64	65.28	97.92	130.56	261.12	652.80	1305.60	13056	6.00	6.00	6.00
4-6 to 5-0 . . . . (10/1/79)	33.62	67.24	100.86	134.48	268.96	672.40	1344.80	13448	6.00	6.01	6.00
5-0 to 5-6 . . . . (4/1/80)	34.63	69.26	103.89	138.52	277.04	692.60	1385.20	13852	6.00	6.01	6.00
5-6 to 6-0 . . . . (10/1/80)	35.67	71.34	107.01	142.68	285.36	713.40	1426.80	14268	6.00	6.00	6.00
6-0 to 6-6 . . . . (4/1/81)	36.74	73.48	110.22	146.96	293.92	734.80	1469.60	14696	6.00	5.99	6.00
6-6 to 7-0 . . . . (10/1/81)	37.84	75.68	113.52	151.36	302.72	756.80	1513.60	15136	6.00	6.03	6.00
7-0 to 7-6 . . . . (4/1/82)	38.98	77.96	116.94	155.92	311.84	779.60	1559.20	15592	6.00	6.00	6.00
7-6 to 8-0 . . . . (10/1/82)	40.15	80.30	120.45	160.60	321.20	803.00	1606.00	16060	6.00	5.98	6.00
8-0 to 8-6 . . . . (4/1/83)	41.35	82.70	124.05	165.40	330.80	827.00	1654.00	16540	6.00	6.00	6.00
8-6 to 9-0 . . . . (10/1/83)	42.59	85.18	127.77	170.36	340.72	851.80	1703.60	17036	6.00	6.01	6.00
9-0 to 9-6 . . . . (4/1/84)	43.87	87.74	131.61	175.48	350.96	877.40	1754.80	17548	6.00	6.02	6.00
9-6 to10-0 . . . . (10/1/84)	45.19	90.38	135.57	180.76	361.52	903.80	1807.60	18076	6.00	5.97	5.97
10-0 2/ . . . . . (4/1/85)	46.54	93.08	139.62	186.16	372.32	930.80	1861.60	18616	6.00 3/	——	——

1/ Month, day, and year on which issues of June 1, 1969, enter each period. For subsequent issue months add the appropriate number of months.
2/ Extended maturity value reached at 15 years 10 months after issue.
3/ Yield on purchase price from issue date to extended maturity date is 5.83 percent.

* For earlier redemption values and yields see appropriate table in Department Circular 653, 9th Revision, as amended and supplemented.
** This table does not apply if the prevailing rate for Series K bonds being issued at the time the extension begins is different from 6.00 percent.

[FR Doc.75-12 Filed 1-2-75;8:45 am]

Title 9—Animals and Animal Products

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

SUBCHAPTER C—INTERSTATE TRANSPORTATION OF ANIMALS (INCLUDING POULTRY) AND ANIMAL PRODUCTS

PART 73—SCABIES IN CATTLE

Area Quarantined

This amendment quarantines a portion of Moore County in Texas because of the existence of cattle scabies. The restrictions pertaining to the interstate movement of cattle from quarantined areas as contained in 9 CFR Part 73, as amended, will apply to the area quarantined.

Accordingly, Part 73, Title 9, Code of Federal Regulations, as amended, restricting the interstate movement of cattle because of scabies is hereby amended as follows:

In § 73.1a, paragraph (a) relating to the State of Texas is amended to read:

§ 73.1a Notice of quarantine.

(a) Notice is hereby given that cattle in certain portions of the State of Texas are affected with scabies, a contagious, infectious, and communicable disease; and, therefore, the following areas in such State are hereby quarantined because of said disease:

(1) That portion of Cochran County comprised of Greer County school land league 85–6, secs. 3, 4, 21 and 22.

(2) That portion of El Paso County comprised of Block #2, Track 8–B of the San Elizario Grant.

(3) That portion of Moore County comprised of sections 321 and 322, Block 44, H & TC Railway Survey.

* * * * *

(Sec. 4–7, 23 Stat. 32, as amended; secs. 1 and 2, 32 Stat. 791–792, as amended; secs. 1–4, 33 Stat. 1264, 1265, as amended; secs. 3 and 11, 76 Stat. 130, 132; 21 U.S.C. 111–113, 115, 117, 120, 121, 123–126, 134b, 134f; 37 FR 28464, 28477; 38 FR 19141.)

Effective date. The foregoing amendment shall become effective December 30, 1974.

The amendment imposes certain further restrictions necessary to prevent the interstate spread of cattle scabies and must be made effective immediately to accomplish its purpose in the public interest. It does not appear that public participation in this rulemaking proceeding would make additional relevant information available to the Department.

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

Done at Washington, D.C., this 30th day of December 1974.

J. M. HEJL,
Deputy Administrator, Veterinary Services, Animal and Plant Health Inspection Service

[FR Doc.75–100 Filed 1–2–75;8:45 am]

PART 73—SCABIES IN CATTLE

Release of Areas Quarantined

These amendments release portions of Stevens County and a portion of Seward County in Kansas and a portion of Cimarron County in Oklahoma from the areas quarantined because of cattle scabies. Therefore, the restrictions pertaining to the interstate movement of cattle from quarantined areas contained in 9 CFR Part 73, as amended, will not apply to the excluded areas, but the restrictions pertaining to the interstate movement of cattle from nonquarantined areas contained in said Part 73 will apply to the excluded areas. No areas in Kansas or Oklahoma remain under quarantine.

Accordingly, Part 73, Title 9, Code of Federal Regulations, as amended, restricting the interstate movement of cattle because of scabies is hereby amended as follows:

In § 73.1a, paragraph (c) relating to the State of Oklahoma and paragraph (d) relating to the State of Kansas are deleted.

(Secs. 4–7, 23 Stat. 32, as amended; secs. 1 and 2, 32 Stat. 791–792, as amended; secs. 1–4, 33 Stat. 1264, 1265, as amended; secs. 3 and 11, 76 Stat. 130, 132; 21 U.S.C. 111–113, 115, 117, 120, 121, 123–126, 134b, 134f; 37 FR 28464, 28477; 38 FR 19141)

Effective date. The foregoing amendments shall become effective on December 30, 1974.

The amendments relieve restrictions no longer deemed necessary to prevent the spread of cattle scabies and should be made effective promptly in order to be of maximum benefit to affected persons. It does not appear that public participation in this rulemaking proceeding would make additional relevant information available to the Department.

Accordingly, under the administrative procedure provisions in 5 U.S.C. 553, it is found upon good cause that notice and other public procedure with respect to 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.

Done at Washington, D.C., this 30th day of December 1974.

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

[FR Doc.75–154 Filed 1–2–75;8:45 am]

SUBCHAPTER D—EXPORTATION AND IMPORTATION OF ANIMALS (INCLUDING POULTRY) AND ANIMAL PRODUCTS

PART 97—OVERTIME SERVICES RELATING TO IMPORTS AND EXPORTS

Commuted Traveltime Allowances

The purpose of this amendment is to establish commuted traveltime periods as nearly as may be practicable to cover the time necessarily spent in reporting to and returning from the place at which an employee of Veterinary Services performs overtime or holiday duty when such travel is performed solely on account of overtime or holiday duty. Such establishment depends upon facts within the knowledge of the Animal and Plant Health Inspection Service.

Therefore, pursuant to the authority conferred upon the Deputy Administrator, Veterinary Services, Animal and Plant Health Inspection Service by § 97.1 of the regulations concerning overtime services relating to imports and exports (9 CFR 97.1), administrative instructions 9 CFR 97.2 (1974 ed.), as amended November 27, 1974 (39 FR 41356–41358), and December 11, 1974 (39 FR 43294), prescribing the commuted traveltime that shall be included in each period of overtime or holiday duty, is hereby amended by adding to or deleting from the respective list therein as follows:

WITHIN METROPOLITAN AREA

ONE HOUR

Add: Lincoln Airport, Lincoln, Nebraska.

(64 Stat. 561; 7 U.S.C. 2260.)

Effective date. The foregoing amendment shall become effective January 3, 1975.

It is to the benefit of the public that this instruction be made effective at the earliest practicable date. It does not appear that public participation in this rulemaking proceeding would make additional relevant information available to the Department.

Accordingly, pursuant to 5 U.S.C. 553, it is found upon good cause that notice and public procedure on this instruction are impracticable, unnecessary, and contrary to the public interest and good cause is found for making it effective less than 30 days after publication in the FEDERAL REGISTER.

Done at Washington, D.C., this 30th day of December 1974.

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

[FR Doc.75–155 Filed 1–2–75;8:45 am]

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

PART 113—STANDARD REQUIREMENTS

Correction and Clarification; Miscellaneous Amendments

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

Section 113.2 is amended by correcting the spelling of "permittee" in the lead paragraph and is further amended by deleting the words "official required" as being unnecessarily restrictive in § 113.2 (a).

The lead paragraph in § 113.3(a) is amended by deleting the words "paragraph (b) of" and by adding a new subparagraph (3) to clarify the requirements for bulk samples prescribed in

§ 113.3(a)(1)(f). Section 113.3(b)(8) is amended to restrict the submission of prelicensing samples to those requested to prevent unnecessary submissions.

Section 113.26 is corrected by changing the spelling of "biological" in the lead paragraph and adding cell lines and primary cells.

Section 113.51 is corrected by changing "of" to "or" in the first sentence of paragraph (d) and deleting the words "fluorescent antibody" in paragraph (e) as being unnecessarily restrictive.

Section 113.65 is corrected by changing the spelling of "peptone" in § 113.65 (b) (1). Section 113.92 is corrected by changing the spelling of "injected" in § 113.92 (c) (2).

Section 113.95 is clarified by inserting the words "shall be used" in § 113.95(c). Section 113.95 (c) (1) is corrected by changing "bled" to "available" in subtoneally" and by changing "MLD_{50} division (c) (3) (iii) of each.

Section 113.96 and § 113.97 are corrected by changing word "have" to "has" in the lead paragraph of each and by changing "bled" to 'available" in subdivision (c) (3) (iii) of each.

Section 113.97 is further corrected by changing the word "at" to "and" in solution." The spelling of "doses" is corperiod at the end of subparagraph (c) (2). Subdivision (c) (4) (vi) is clarified by inserting the words "one ml of this solution." The spelling of "doses" is corrected in subdivision (c) (5) (iii).

Sections 113.101, 113.102, and 113.103 are amended by changing the names of the biological products affected for scientific accuracy. The phrase "Avian Isolates" is substituted for the word "avicida" in the caption and lead paragraph of §§ 113.101, 113.102, and 113.103. In addition, § 113.102 is further changed by inserting the word "be" in subparagraph (c) (3) as an editorial correction.

The severity of the challenge is adjusted in § 113.104(d)(3) by reducing the dose of challenge culture from 0.5 ml to 0.2 ml as being more realistic and scientifically correct. Subparagraph 113.104(d)(5) is reworded to clarify the dilutions to be used.

The lead paragraph in § 113.251(a) is worded for clarity. Section 113.123 is corrected for scientific accuracy and to conform to §113.139 by substituting "Feline Panleukopenia" for "feline distemper" in each place it appears.

Section 113.201(e) is amended for scientific accuracy. § 113.202(a) is amended by increasing the acceptable range in the packed cell requirements as being more realistic and correct. The name of Mycoplasma Gallisepticum Plate Antigen is corrected in paragraph § 113.202(d).

The lead paragraph in § 113.251 (a) is amended to recognize the difference in accuracy for cylinders customarily used for measuring small and large volumes. § 113.251 is further amended to relax the guinea pig size requirements in subparagraph (d) (2) by proving a weight range of 340 to 380 grams instead of a specified 350 gram weight.

Section 113.252(c)(2) is corrected by deleting the words "do not." Section 113.255(c)(3)(iii) is amended by correcting the spelling of "antitoxin."

1. The lead paragraph in § 113.2 and the provisions in paragraph (a) are revised to read:

§ 113.2 Testing aids.

To better insure consistent and reproducible test results when Standard Requirement tests prescribed in the regulations are conducted, Veterinary Services Laboratories, U.S. Department of Agriculture, may provide testing aids, when available, to licensees, permittees, and applicants for licenses and permits. Such aids shall be as follows:

(a) Supplemental Assay Method (SAM) is a technical bulletin containing detailed instructions for conducting a test. Such instructions shall be in accordance with the procedures currently being followed at Veterinary Services Laboratories and as improved, proven procedures are developed, shall be revised and reissued prior to application.

* * * * *

2. The lead paragraph in § 113.3(a) and the provisions in subparagraph (a) (3), and subparagraph (b)(8) are amended to read:

§ 113.3 Sampling of biological products.

* * * * *

(a) An employee of the Department, of the licensee, or of the permittee, as designated by the Deputy Administrator shall select prerelease samples of biological product to be tested by Veterinary Services. Such samples shall be forwarded to the place designated by the Deputy Administrator and in the number prescribed in this section.

* * * * *

(3) When bulk samples of completed product in liquid form are to be tested as prescribed in subparagraph (1) of this paragraph, the number of such samples from each serial and the minimum quantity of product to be provided in each sample shall be stated in the filed Outline of Production.

(b) * * *

(8) *Prelicensing.* Samples for prelicensing of biological product shall be submitted upon request from Veterinary Services. Such samples shall be double the number prescribed in this section for such product.

* * * * *

3. The lead paragraph in § 113.26 is amended to read:

§ 113.26 Detection of viable bacteria and fungi except in live vaccine.

Each serial and subserial of biological product except live vaccines shall be tested as prescribed in this section unless otherwise specified by the Deputy Administrator. When cell lines, primary cells, or ingredients of animal origin used in the preparation of a biological product are required to be free of viable bacteria and fungi, they shall also be tested as prescribed in this section.

* * * * *

4. The introductory portion of paragraph (d) in § 113.51 and the provisions in paragraph (e) are amended to read:

§ 113.51 Requirements for primary cells used in biological product production.

* * * * *

(d) Each batch of primary cells of bovine origin or each subculture of such cells used to prepare a biological product shall be shown free of Bovine Virus Diarrhea (BVD) virus. The samples for testing shall consist of at least 10 monolayers of cells, each with an area at least as large as a 10.5 × 22 mm coverslip. The samples for testing shall be obtained from at least the second subpassage from intact tissue. The monolayers shall be grown to at least 80 percent confluency using the media (with additives) intended for growth and maintenance and under conditions similar to those used to prepare the product. At least five of the monolayers shall be inoculated with BVD virus as positive controls. All monolayers shall be further incubated at 35–37° C for an additional 4 to 6 days. All monolayers shall then be removed from their media, processed, and stained with anti-BVD fluorescein-tagged antibody conjugate, and examined for presence of specific fluorescence attributable to BVD virus.

* * * * *

(e) Each batch of primary cells or each subculture of cells used to prepare a biological product shall be shown free of other specific viruses using applicable tests.

* * * * *

5. § 113.65(b)(1) is amended to read:

§ 113.65 Brucella Abortus Vaccine.

* * * * *

(b) *Potency test.* * * *

(1) At least four single-dose or two multiple-dose final container samples of completed product shall be tested for the number of viable organisms per cubic centimeter of rehydrated vaccine. A bacterial count shall be made on tryptose agar plates from suitable dilutions using 1 percent peptone as a diluent.

* * * * *

6. § 113.92 (c) (2) is amended to read:

§ 113.92 Clostridium Hemolyticum Bacterin.

* * * * *

(c) *Potency test.* * * *

(2) Clostridium hemolyticum challenge material, available upon request from Veterinary Services, shall be used for challenge 14 to 15 days following the last injection of the product. Each of the eight vaccinates and each of five additional nonvaccinated guinea pigs for controls shall be injected intramuscularly with approximately 100 LD_{50} of challenge material. This dose shall be

determined by statistical analysis of results of titrations of the challenge material. The vaccinates and controls shall be observed for 3 days post-challenge and all deaths recorded.

* * * *

7. Sections 113.95 (c) and (c)(1) are revised to read:

§ 113.95 Clostridium Botulinum Type C Bacterin-Toxoid.

* * * * *

(c) *Potency test.* Bulk or final container samples of completed product from each serial shall be tested for potency, using susceptible mink as test animals. At least five vaccinates and three unvaccinated controls of the same source and approximately the same age shall be used.

(1) Each of the vaccinates shall be injected subcutaneously with the dose recommended on the label for mink. Twenty-one to twenty-eight days post-injection, the vaccinates and the controls shall be challenged intraperitoneally with botulinum Type C toxin which has been titrated in mice to provide for a $10^{4.6}$ mouse MLD dose. The titration technique shall include inoculation of the mice intraperitoneally.

* * * * *

8. The introductory text in § 113.96 and the provisions in paragraph (c)(3)(iii) are revised 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.

* * * * *

(c) *Potency test.* * * *

(3) * * *

(iii) If less than four rabbits are available, the test is invalid and shall be repeated; *Provided*, That, if the test is not repeated, the serial shall be declared unsatisfactory.

* * * * *

9. The introductory text in § 113.97, and the provisions in paragraphs (c)(1)(vi), (c)(2), (c)(3)(iii), (c)(4)(iv), and (c)(5)(iii) are revised 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) *Potency test.* * * *

(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 grams 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.

(2) Each of at least eight rabbits, each weighing 4–8 pounds, shall be injected subcutaneously with not more than half of the recommended sheep dose. The dose for a combination product having both Type C and Type D fractions shall be half of the recommended cattle dose; *Provided*, That, if the product is recommended only for sheep, half of the recommended sheep dose shall be used. A second dose shall be given not less than 20 days nor more than 23 days after the first dose.

(3) * * *

(iii) If less than four rabbits are available, the test is invalid and shall be repeated; *Provided*, That, if the test is not repeated, the serial shall be declared unsatisfactory.

(4) * * *

(iv) Dilute 1 ml of serum with 1 ml of diluent (1:2) and combine 1 ml of this solution with 10 L_+ doses of diluted Standard Toxin.

(5) * * *

(iii) If any mice inoculated with the mixture of serum with 10 L_+ doses of Standard Toxin die, the serum is considered to contain less than 2 International Units per ml.

* * * * *

10. The heading and introductory text in § 113.101 are revised to read:

§ 113.101 General Requirements for Pasteurella Multocida Bacterins, Avian Isolates.

Pasteurella Multocide Bacterin, Avian Isolates, shall be prepared with cultures of *Pasteurella multocida*, avian isolates, Type 1 or Type 3 or both (Little and Lyons Classification) which have been inactivated and are nontoxic; *Provided*, That, avian isolates other than Types 1 and 3 may be added if written into the filed Outline of Production for the product.

* * * * *

11. The heading and introductory text in § 113.102 and the provisions in paragraph (c)(3) are revised to read:

§ 113.102 Pasteurella Multocida Bacterin, Avian Isolates, Type 1.

Each serial of Pasteurella Multocida Bacterin, Avian Isolates, prepared with Type 1 strains, shall be tested as prescribed in this section. A serial found unsatisfactory by any prescribed test shall not be released.

* * * * *

(c) *Potency test.* * * *

(3) *Unvaccinated controls.* Each of not more than 21 chickens shall be held as controls.

* * * * *

12. The heading and introductory text in § 113.103 are revised to read:

§ 113.103 Pasteurella Multocida Bacterin, Avian Isolates, Type 3.

Each serial of Pasteurella Multocida Bacterin, Avian Isolates, prepared with Type 3 strains, shall be tested as prescribed in this section. A serial found unsatisfactory by any prescribed test shall not be released.

* * * * *

13. Section 113.104 (d)(3), and (d)(5), are revised to read:

§ 113.104 Erysipelas Bacterin.

* * * * *

(d) * * *

(3) Each injected mouse shall be challenged subcutaneously 14–21 days after being injected with the diluted bacterin. A 0.2 ml dose containing at least 100 mouse LD_{50} of a suitable culture of *Erysipelothrix insidiosa* shall be used. All survivors in each group of mice shall be recorded 10 days post-challenge.

* * * * *

(5) Using the same three consecutive dilutions of the Standard and Unknown, obtain the total survivors of each. If the total number of survivors for the Standard exceeds the total number of survivors for the Unknown by a number greater than six, the Unknown is unsatisfactory.

* * * * *

14. The introductory text in § 113.121 is revised to read:

§ 113.121 Canine Distemper Vaccine, Killed Virus.

Canine Distemper Vaccine, Killed Virus, shall be prepared from virus-bearing cell culture fluids or tissues obtained from animals that have developed canine distemper following inoculation with virulent canine distemper virus. Each serial shall meet the applicable general requirements prescribed in § 113.120 and special requirements prescribed in this section. Any serial found unsatisfactory by a prescribed test shall not be released.

* * * * *

15. The heading and introductory text in § 113.123, the introductory text of paragraph (b) and the provisions in subparagraphs (b)(2) and (3) are revised to read:

§ 113.123 Feline Panleukopenia Vaccine, Killed Virus.

Feline Panleukopenia Vaccine, Killed Virus, shall be prepared from virus-bearing cell culture fluids or from tissues obtained from cats that have developed feline panleukopenia following inoculation with virulent feline panleukopenia virus. Each serial shall meet the

applicable requirements prescribed in § 113.120 and special requirements prescribed in this section. Any serial found unsatisfactory by a prescribed test shall not be released.

* * * * *

(b) *Potency test.* Bulk or final container samples of completed product shall be tested for potency using four feline panleukopenia susceptible cats (two vaccinates and two controls). The susceptibility of the cats shall be determined by a constant virus-varying serum neutralization test in tissue culture using 100 to 300 $TCID_{50}$ of virus. Susceptible cats shall have no neutralization at a 1:2 serum dilution.

* * * * *

(2) *Challenge.* At the end of the postvaccination observation period, the two vaccinates and the two controls shall be exposed to virulent feline panleukopenia virus and observed each day for an additional 14 days. White blood cell counts shall be made on the vaccinates and the controls for 9 consecutive days following challenge.

(3) *Interpretation.* If the control cats do not develop signs of feline panleukopenia including pronounced leukopenia, wherein the white cell count drops to 4,000 or less per cubic mm within the test period or the white cell drops to less than 25 percent of the normal level established by an average of three or more counts taken prior to the onset of leukopenia, the test is inconclusive and may be repeated; *Provided*, That, if the vaccinates show a pronounced leukopenia or do not remain free of feline panleukopenia, the serial is unsatisfactory.

16. Section 113.201 (e) is revised to read:

§ 113.201 Pullorum antigen.

* * * * *

(e) *Homogeneity requirement.* Antigens shall show no evidence of autoagglutination or unusual appearance such as the presence of flakes, specks, or a preponderance of filament forms. Microscopic examination shall be made in this determination.

* * * * *

17. Section 113.202 (a) and (d) are revised to read:

§ 113.202 Avian Mycoplasma Antigen.

* * * * *

(a) *Density requirements.* A 2.5 ml sample of completed antigen shall be diluted with 2.5 ml of Sorenson's buffer solution (use buffer solution at pH 6.0 for Mycoplasma Gallisepticum Plate Antigen and at pH 7.0 for Mycoplasma Gallisepticum Tube Antigen and Mycoplasma Synoviae Plate Antigen) in a modified Hopkins tube and sedimented at 1,000 x g in a refrigerated certifuge at 20° C for 90 minutes. If the packed cell volume of the completed antigen is not 1.2 percent (± 0.4 percent), the serial is unsatisfactory.

* * * * *

(d) *Hydrogen ion concentration.* The hydrogen ion concentration shall be determined with a pH meter which has been standardized with a pH buffer just prior to use. The pH of Mycoplasma Gallisepticum Plate Antigen shall be 6.0 ± 0.2; the pH of Mycoplasma Gallisepticum Tube Antigen and Mycoplasma Synoviae Plate Antigen shall be 7.0 ± 0.2.

* * * * *

18. The introductory text in § 113.251 (a) and the provisions in paragraph (d) (2) are revised to read:

§ 113.251 Tetanus Antitoxin.

* * * * *

(a) *General requirements.* The amount of antitoxin in a final container shall be the amount which is delivered from such container when opened and inverted until the flow stops. A graduated volumetric cylinder which conforms to the National Bureau of Standards requirements shall be used. The reading shall be made at the bottom of the meniscus. Volumes of 10 ml or less shall be recorded to the nearest 0.1 and volumes over 10 ml shall be recorded to the nearest ml.

* * * * *

(d) *Potency test.* * * *

(2) The standard toxin test dose is that amount which when mixed with 0.1 unit of standard antitoxin, incubated at 20 to 25° C for 1 hour, and injected subcutaneously into a 340 to 380 gram guinea pig, results in death of that guinea pig within approximately 96 hours with clinical signs of tetanus. The toxin shall be diluted so the test dose shall be in 2.0 ml.

19. Section 113.252(c)(2) is revised to read:

§ 113.252 Swine Erysipelas Antiserum.

* * * * *

(c) *Potency test.* * * *

(2) If less than eight of the 10 controls die from erysipelas within 7 days postchallenge, the test is invalid. All dead mice shall be examined to determine if the cause of death was *Erysipelothrix insidiosa* infection.

* * * * *

20. Section 113.255(c)(3)(iii) is revised to read:

§ 113.255 Clostridium Perfringens Type D Antitoxin.

* * * * *

(c) *Potency test.* * * *

(3) * * *

(iii) If any mice inoculated with the mixture of Clostridium Perfringens Type D Antitoxin diluted 1:34 and 10 L_o doses of Standard Toxin die, the antitoxin is considered to contain less than 34 International Units per ml and the serial is unsatisfactory.

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

These amendments make editorial changes to correct printing, grammar, and spelling errors, to relax requirements where indicated, and to clarify questionable regulations without making other substantive changes. 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, D.C., this 30th day of December 1974.

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

[FR Doc.75-156 Filed 1-2-75;8:45 am]

Title 16—Commercial Practices

CHAPTER I—FEDERAL TRADE COMMISSION

SUBCHAPTER A—PROCEDURES AND RULES OF PRACTICE

DEPUTY BUREAU DIRECTORS

Authority; Miscellaneous Amendments

The Commission announces the following amendments to Chapter I of Title 16 of the Code of Federal Regulations to give Deputy Bureau Directors the same authority conferred upon Assistant Bureau Directors. The amendments are effective on January 3, 1975.

PART 2—NONADJUDICATIVE PROCEDURES

Subpart A—Investigations

Section 2.1 is revised to read as follows:

§ 2.1 How initiated.

Commission investigations and inquiries may be originated upon the request of the President, Congress, governmental agencies, or the Attorney General; upon referrals by the courts; upon complaint by members of the public; or by the Commission upon its own initiative. The Commission has delegated to the Directors, Deputy Directors, and Assistant Directors of the Bureaus of Competition and Consumer Protection, and the Regional Directors and Assistant Regional Directors of the Commission's regional offices, without power of redelegation, limited authority to initiate investigations.

Section 2.7 is revised to read as follows:

§ 2.7 Subpoenas in investigations.

(a) The Commission or any member thereof may issue a subpoena, directing the person named therein to appear before a designated representative at a designated time and place to testify or to produce documentary evidence, or both, relating to any matter under investigation by the Commission. The Directors, Deputy Directors, and Assistant Directors of the Bureaus of Competition, Consumer Protection, and Economics, and the Regional Directors and Assistant Regional Directors of the Commission's regional offices, pursuant to delegation of authority by the Commission, without power of redelegation, also may issue investigational subpoenas, and, for good cause shown, may extend the time prescribed

for compliance with subpoenas issued during the investigation of any matter. The Director, Deputy Director, Assistant Director, Regional Director, or Assistant Regional Director, who issues any subpoena under this section is authorized to negotiate and approve the terms of satisfactory compliance therewith.

(b) Any motion to limit or quash any investigational subpoena shall be filed with the Secretary of the Commission, within ten (10) days after service of the subpoena, or, if the return date is less than ten (10) days after service of the subpoena, within such other time as may be allowed. All motions to limit or quash any investigational subpoenas shall be ruled upon by the Commission itself, but the above-designated Directors, Deputy Directors, Assistant Directors, Regional Directors and Assistant Regional Directors are delegated, without power of redelegation, the authority to rule upon motions for extensions of time within which to file motions to limit or quash any investigational subpoenas.

Section 2.11 is revised to read as follows:

§ 2.11 Orders requiring access.

(a) The Commission may issue an order requiring any corporation being investigated to grant access to files for the purpose of examination and the right to copy any documentary evidence. The Directors, Deputy Directors, and Assistant Directors of the Bureaus of Competition, Consumer Protection, and Economics and the Regional Directors and Assistant Regional Directors of the Commission's regional offices, pursuant to delegation of authority by the Commission, without power of redelegation, are authorized, for good cause shown, to extend the time prescribed for compliance with orders requiring access issued during the investigation of any matter.

(b) Any motion to limit or quash an order requiring access shall be filed with the Secretary of the Commission within ten (10) days after service of the order, or, if the date for compliance is less than ten (10) days after service of the order, within such other time as may be allowed. All motions to limit or quash orders requiring access shall be ruled upon by the Commission itself, but the above-designated Directors, Deputy Directors, Assistant Directors, Regional Directors and Assistant Regional Directors are delegated, without power of redelegation, the authority to rule upon motions for extensions of time within which to file motions to limit or quash orders requiring access.

Section 2.12 is revised to read as follows:

§ 2.12 Reports.

(a) The Commission may issue an order requiring a corporation to file a report or answers in writing to specific questions relating to any matter under investigation. The Directors, Deputy Directors, and Assistant Directors of the Bureaus of Competition, Consumer Protection, and Economics, and the Regional Directors and Assistant Regional Directors of the Commission's regional offices, pursuant to delegation of authority by the Commission, without power of redelegation, are authorized, for good cause shown, to extend the time prescribed for compliance with orders requiring reports or answers to questions issued during the investigation of any matter.

(b) Any motion to limit or quash an order requiring a report or answers to specific questions shall be filed with the Secretary of the Commission within ten (10) days after service of the order, or, if the date for compliance is less than ten (10) days after service of the order, within such other time as may be allowed. All motions to limit or quash orders requiring reports or answers to questions shall be ruled upon by the Commission itself, but the above-designated Directors, Deputy Directors, Assistant Directors, Regional Directors and Assistant Regional Directors are delegated, without power of redelegation, the authority to rule upon motions for extensions of time within which to file motions to limit or quash orders requiring reports or answers to questions.

Section 2.14(c) is revised to read as follows:

§ 2.14 Disposition.

* * * * *

(c) The Commission has delegated to the Directors, Deputy Directors, and Assistant Directors of the Bureaus of Competition and Consumer Protection, without power of redelegation, limited authority to close investigations. The closing action of a Bureau Director, Deputy Bureau Director or Assistant Bureau Director does not become effective until the files have been sent to the Secretary of the Commission and no member of the Commission has objected within five (5) working days after receiving the notice to close from the Secretary.

PART 3—RULES OF PRACTICE FOR ADJUDICATIVE PROCEEDINGS

Subpart G—Reports of Compliance

Section 3.61(c) is revised to read as follows:

§ 3.61 Reports of compliance.

* * * * *

(c) The Commission has delegated to the Directors, Deputy Directors, and Assistant Directors of the Bureaus of Competition and Consumer Protection, without power of redelegation, the authority for good cause shown, to extend the time within which reports of compliance with orders to cease and desist may be filed. It is to be noted, however, that an extension of time within which a report of compliance may be filed, or the filing of a report which does not evidence full compliance with the order, does not in any circumstances suspend or relieve a respondent from his obligation under the law with respect to compliance with such order. An order of the Commission to cease and desist becomes final on the date and under the conditions provided in section 5 (g), (h), (l), (j), and (k) of the Federal Trade Commission Act (15 U.S.C. 45 (g), (h), (l), (j), and (k)) and section 11 (g), (h), (l), (j), and (k) of an Act to supplement existing laws against unlawful restraints and monopolies, and for other purposes, as amended—the Clayton Act, as amended (15 U.S.C. 21 (g), (h), (l), (j), and (k)). Any person, partnership or corporation against which an order to cease and desist has been issued who is not in full compliance with such order on and after the date provided in these statutes for the order to become final is in violation of such order and is subject to an immediate action for civil penalties.

* * * * *

PART 4—MISCELLANEOUS RULES

Section 4.2(a) is revised to read as follows:

§ 4.2 Requirements as to form and filing of documents other than correspondence.

(a) *Filing.*—Except as otherwise provided, all documents submitted to the Commission shall be addressed to and filed with the Secretary of the Commission: *Provided, however,* That in any instance informal applications or requests may be submitted directly to the official in charge of any office of the Commission or to the Director, Deputy Director, or Assistant Director of the appropriate bureau or office.

* * * * *

(Sec. 6, 38 Stat. 721 (15 U.S.C. 46).)

By direction of the Commission dated December 19, 1974.

[SEAL] CHARLES A. TOBIN, *Secretary.*

[FR Doc.75-135 Filed 1-2-75;8:45 am]

PART 13—PROHIBITED TRADE PRACTICES AND AFFIRMATIVE CORRECTIVE ACTIONS

Subpart—Corrective Actions and/or Requirements

The Federal Trade Commission announces the following amendments to Part 13, Subchapter A of Chapter I of Title 16 to change the title of Part 13 and establish a new subpart providing for corrective actions and/or requirements.

The title of Part 13 is changed from "Prohibited Trade Practices" to "Prohibited Trade Practices and Affirmative Corrective Actions."

The following new subpart and codification is added following Subpart—Controlling, Unfairly, Seller-Suppliers:

Subpart—Corrective Actions and/or Requirements

§ 13.533 Corrective actions and/or requirements.
§ 13.533-5 Arbitration.
§ 13.533-10 Corrective advertising.
§ 13.533-15 Destruction of records and/or data.
§ 13.533-20 Disclosures.
§ 13.533-25 Displays, in-house.
§ 13.533-30 Election of officers.

AUTHORITY: Sec. 6(g), 5, 38 Stat. 722, 719 (15 U.S.C. 46, 45); sec. (a)(1), 80 Stat. 383 (5 U.S.C. 552).

By direction of the Commission, dated December 26, 1974.

[SEAL] CHARLES A. TOBIN, *Secretary.*

[FR Doc.75-134 Filed 1-2-75;8:45 am]

Title 36—Parks, Forests and Memorials

CHAPTER I—NATIONAL PARK SERVICE, DEPARTMENT OF THE INTERIOR

PART 7—SPECIAL REGULATIONS, AREAS OF THE NATIONAL PARK SYSTEM

Lake Meredith Recreation Area; Off Road Use

A proposal was published at page 17851 of the FEDERAL REGISTER of May 21, 1974, to change the title for § 7.57 now reading Sanford Recreation Area to Lake Meredith Recreation Area and to revise paragraph (a) now designated as "Reserved" to establish areas for use by off-road vehicles. Interested persons were given thirty days within which to submit written comments, suggestions, or objections with respect to the proposed amendment. No comments, suggestions, or objections have been received and the proposed amendments are hereby adopted without change and are set forth below. These amendments shall take effect February 3, 1975.

The heading for § 7.57, *Sanford Recreation Area*, is revised to read § 7.57 *Lake Meredith Recreation Area* and paragraph (a) now designated as reserved is added to read as follows:

§ 7.57 Lake Meredith Recreation Area.

(a) The operation of motor vehicles within the Lake Meredith Recreation Area is prohibited outside of established public roads, parking areas, except within the cutbanks of Blue Creek, comprising about 275 acres, and except below the 3,000 ft. contour on the following described lands, being known as the Rosita Area on the Canadian River flood plain:

(1) Beginning at property corner 191 at coordinates 536,112.90N and 1,894,-857.49E thence in a straight line S05°14'47" E, 3349.09 ft. to property corner 192, thence in a straight line N85°03'12" E, 6999.38 ft., to property corner 193, thence in a straight line N58°29'53" E, 3737.77 ft., to property corner 194, thence in a straight line N51°20'25" E, 1457.45 ft., to property corner 195, thence in a straight line S74°40'44" E, 4064.61 ft., to property corner 196, thence in a straight line N79°59'22" E, 3118.40 ft. to property corner 197A, thence in a northeasterly direction to property corner 200, thence in a straight line N56°24'11" E, 1073.57 ft., to property corner 201, thence in a straight line S80°04'22" E, 2684.69 ft., to property corner 202, thence in a straight line N69°21'31" E, 2974.09 ft. to property corner 203, thence in a straight line S37°59'16" E, 1538.83 ft., to property corner 204, thence in a straight line N28°36'59" E, 744.10 ft., to property corner 205, thence in a straight line N00°19'04" E, 1136.41 ft., to property corner 206, thence in a westerly direction to property corner 181, thence in a straight line S89°51'52" W, 1434.80 ft. to property corner 182, thence in a straight line N75°53'25" W, 4267.11 ft., to property corner 183, thence in a straight line S76°16'20" W, 3835.45 ft., to property corner 184, thence in a westerly direction to property corner 189, thence in a straight line S71°35'59" W, 2901.46 ft., to property corner 190, thence in a straight line S78°24'18" W, 6506.70 ft. to the point of beginning as shown on Bureau of Reclamation drawing number 662-525-1431 dated July 9, 1965, such Rosita Area comprising about 1,500 acres.

(2) Nothing contained in this § 7.57 (a) shall be deemed to restrict the use of motor vehicles outside of public roads and parking areas for official or emergency purposes, as required in the discretion of the Superintendent.

(3) The Superintendent may establish limits on the number of vehicles permitted in the above designated areas when such limitations are necessary in the interest of public safety or for coordination of other visitor uses, or for conservation of the natural resources of the area.

WILLIAM E. DYER, *Superintendent.*

[FR Doc.75-136 Filed 1-2-75;8:45 am]

Title 50—Wildlife and Fisheries

CHAPTER I—U.S. FISH AND WILDLIFE SERVICE, DEPARTMENT OF THE INTERIOR

PART 28—PUBLIC ACCESS, USE, AND RECREATION

Wichita Mountains Wildlife Refuge, Okla.

The following special regulations are issued and are effective January 3, 1975.

§ 28.28 Special regulations; public access, use, and recreation; for individual wildlife refuge areas.

OKLAHOMA

WICHITA MOUNTAINS WILDLIFE REFUGE

Those portions of the Wichita Mountains Wildlife Refuge, Oklahoma, designated for public use are open for certain recreational uses from January 1 through December 31, 1975, inclusive. The public use area totals approximately 22,400 acres and is delineated on maps available at refuge headquarters, Cache, Oklahoma, and from the Regional Director, U.S. Fish and Wildlife Service, P.O. Box 1306, Albuquerque, New Mexico 87103. Public access, use and recreational activity shall be in accordance with all Federal and State laws and regulations and all official signs posted in the area subject to the following special conditions:

(1) Sightseeing, nature observation, photography and hiking are permitted.

(2) Camping and picnicking are permitted in recreation areas containing facilities for these purposes unless prohibited by signs. Exceeding posted visiting hours or unit capacities of these areas is prohibited. A written permit is required for stays exceeding seven (7) days.

(3) Fires are permitted only in recreation areas where camping or picnicking is allowed and only at such times or hours that the areas are open to these uses. Dead, fallen timber may be used.

(4) Boating is permitted only on Elmer Thomas Lake. All other floating devices are prohibited on all refuge waters unless permitted by other Federal regulations. Boating is prohibited in marked scuba diving and swimming areas.

(5) Swimming, wading, snorkeling and skin diving are permitted only at designated swimming beaches, and only when these beaches are manned by refuge supervised lifeguards. Lifejackets and buoyant vests may be worn while swimming. Food, beverages and pets are prohibited on swimming beaches. Beach users must comply with all official beach signs posted on the area and with the directions of authorized lifeguards.

(6) Scuba diving is permitted only on Elmer Thomas Lake. Diving areas must be marked with appropriate warning flags when outside of marked swimming areas. Flags must be removed before leaving the area. Inflatable vests may be worn while diving.

(7) Pets must be kept on leash.

(8) Vehicles found parked in any closed area, any "no parking" area, or in any area after posted visiting hours may be removed from the area. Any charges or expenses incurred by such removal, including storage fees, shall be borne by the owner of the vehicle.

(9) The use of gliders, including hanggliders, is prohibited.

(10) Possession or use of any alcoholic beverage by persons under twenty-one (21) years of age is prohibited.

The provisions of this special regulation supplement the regulations which govern access, use, and recreation on wildlife refuge areas generally which are set forth in Title 50, Code of Federal Regulations, Part 28, and are effective through December 31, 1975.

§ 33.5 Special regulations; sport fishing; for individual wildlife refuge areas.

Sport fishing on the Wichita Mountains Wildlife Refuge, Oklahoma, is per-