Calculation of the allocation of the shortfall due to sovereign action—Germany ($200 shortfall to be allocated):

Customer	Allocation share	Allocation share of actual shortfall	Actual shortfall allocated
C	$50/$150	33.3% of $200	$66.67
D	$100/$150	66.7% of $200	$133.33
Total			$200.000

This would result in the claims of customers C and D being reduced below zero.

Accordingly, the claims of customer C and D will only be reduced to zero, or $50 for C and $100 for D. This results in a Total Excess Shortfall of $50.

Actual shortfall	Allocation of shortfall for customer C	Allocation of shortfall for customer D	Total excess shortfall
$200	$50	$100	$50

This shortfall will be divided among the remaining customers who have authorized funds to be held outside the U.S. or in a currency other than U.S. dollars.

Customer	Total claims of customers permitting funds to be held outside the U.S.	Portion of claim required to be int the U.S.	Allocation share (column B–C/column B Total—all customer claims in U.S.)	Allocation share of actual total excess shortfall	Actual total excess shortfall allocated
B	$100	$50	$50/$200	25% of $50	$12.50
C	50	0	[1]		0
D	200	100	$100/200	50% of $50	25
E	100	50	50/100	25% of $50	12.50
Total	450.00				50.00

[1] Claim already reduced to $0.

Claims After Reductions

Customer	Claim in U.S. dollars after allocated non-sovereign shortfall	Allocation of shortfall due to sovereign action Germany	Allocation of total excess shortfall	Claim after all reductions
A	$50			$50.00
B	100		12.50	87.50
C	50	50		0
D	200	100	25	75.00
E	100		12.50	87.50
Total	500.00	150.00	50.00	300.00

Issued in Washington, DC on January 29, 2003, by the Commission.

Jean A. Webb,

Secretary of the Commission.

[FR Doc. 03–2508 Filed 2–3–03; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 529

Certain Other Dosage Form New Animal Drugs; Formalin Solution

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; technical amendment.

SUMMARY: The Food and Drug Administration (FDA) is amending the animal drug regulations to reflect approval of a supplemental new animal drug application (NADA) filed by Natchez Animal Supply Co. The supplemental NADA provides for use of formalin in a water bath for the control of certain external parasites on finfish and shrimp and for the control of certain fungi on finfish eggs. Minor corrections to the regulations are also being made.

DATES: This rule is effective February 4, 2003.

FOR FURTHER INFORMATION CONTACT: Joan C. Gotthardt, Center for Veterinary Medicine (HFV–131), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–827–7571, e-mail: *jgotthar@cvm.fda.gov*.

SUPPLEMENTARY INFORMATION: Natchez Animal Supply Co., 201 John R. Junkin Dr., Natchez, MS 39120, filed a supplement to NADA 137–687 that provides for use of formalin in a water bath for the control of certain external parasites on finfish and shrimp and for the control of certain fungi on finfish eggs. The supplemental NADA is approved as of November 25, 2002, and the regulations are amended in 21 CFR 529.1030 to reflect the approval. The basis of approval is discussed in the freedom of information summary.

In accordance with the freedom of information provisions of 21 CFR part

20 and 21 CFR 514.11(e)(2)(ii), a summary of safety and effectiveness data and information submitted to support approval of this application may be seen in the Dockets Management Branch (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852, between 9 a.m. and 4 p.m., Monday through Friday.

The agency has determined under 21 CFR 25.33(a)(1) that this action is of a type that does not individually or cumulatively have a significant effect on the human environment. Therefore, neither an environmental assessment nor an environmental impact statement is required.

This rule does not meet the definition of ''rule'' in 5 U.S.C. 804(3)(A) because it is a rule of ''particular applicability.'' Therefore, it is not subject to the congressional review requirements in 5 U.S.C. 801–808.

List of Subjects in 21 CFR Part 529

Animal drugs.

Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs and redelegated to the Center for Veterinary Medicine, 21 CFR part 529 is amended as follows:

PART 529—CERTAIN OTHER DOSAGE FORM NEW ANIMAL DRUGS

1. The authority citation for 21 CFR part 529 continues to read as follows:

Authority: 21 U.S.C. 360b.

§ 529.1030 [Amended]

2. Section 529.1030 *Formalin solution* is amended as follows:

(a) In the section heading and in paragraph (a) by removing the word ''solution'' following the word ''Formalin'';

(b) By revising the introductory text of paragraph (b);

(c) In paragraph (b)(1) by removing ''No. 050378'' and by adding in its place ''Nos. 049968 and 050378'';

(d) In paragraph (b)(2) by removing ''Nos. 049968 and'' and by adding in its place ''No.'';

(e) In paragraph (d)(2)(i), in the table, in the heading to the second column, by adding ''daily'' after ''1 hour''; and

(f) In paragraph (d)(2)(iv), in the first column in the table by removing ''½F'' each time it occurs and by adding in its place ''°F''.

The revision is to read as follows:

§ 529.1030 Formalin.

* * * * *

(b) *Sponsors.* See sponsors in § 510.600(c) of this chapter for uses as in paragraph (d) of this section.

* * * * *

Dated: January 21, 2003.

Steven D. Vaughn,

Director, Office of New Animal Drug Evaluation, Center for Veterinary Medicine.

[FR Doc. 03–2601 Filed 2–3–03; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF JUSTICE

Bureau of Prisons

28 CFR Part 522

[BOP–1110–I]

RIN 1120–AB08

Admission and Orientation Program: Removal From Rules

AGENCY: Bureau of Prisons, Justice.

ACTION: Interim final rule.

SUMMARY: In this document, the Bureau of Prisons (Bureau) removes its rules on the Admission and Orientation Program from the CFR. We intend this amendment to streamline our regulations by removing internal agency management procedures that need not be stated in regulation.

DATES: This rule is effective February 4, 2003. Please send comments on this rulemaking by April 7, 2003.

ADDRESSES: Rules Unit, Office of General Counsel, Bureau of Prisons, HOLC Room 754, 320 First Street, NW., Washington, DC 20534.

FOR FURTHER INFORMATION CONTACT: Sarah Qureshi, Office of General Counsel, Bureau of Prisons, phone (202) 307–2105.

SUPPLEMENTARY INFORMATION: In this document, the Bureau of Prisons (Bureau) removes its rules on the Admission and Orientation Program by reserving 28 CFR subpart E. Although we are removing these rules from the CFR, they will remain in Bureau policy statements on the Admission and Orientation Program.

Why Are We Making This Change?

We intend this change to streamline our regulations by removing internal agency management procedures that need not be stated in regulation. In doing this, we will be able to adjust our Admission and Orientation program, through policy instead of rules, to allow us to provide more current information more quickly to new inmates. Bureau policy is a more appropriate vehicle through which to provide instruction and guidance to staff.

Admission and Orientation Program Rules

The three rules in 28 CFR subpart E, §§ 522.40, 522.41, and 522.43 contained descriptions of the Bureau's Admission and Orientation Program. Although we are removing these rules from the CFR, we retain the language of these rules in our Admission and Orientation policy, which is an instructional document for Bureau employees and institutional staff.

Section 522.40 required institutions and staff to ''offer each newly committed inmate an orientation to the institution'' which includes information on the inmate's rights, responsibilities, obligations, and the institution's programs and disciplinary system.

Section 522.41 delineated Warden and staff responsibility for conducting the Admission and Orientation (A&O) program. This section required staff involved in the A&O program to develop an outline of information to present during A&O and develop written orientation materials. This section also instructed staff to monitor inmates with significant emotional stress during A&O, so that the institution could provide them with appropriate assistance.

Section 522.42 contained guidelines for institutions' A&O programs, including such details as location, activities, and length of the program.

All of these rules consist of our instruction and guidance to Bureau staff. These rules relate solely to internal agency management and practice, and do not impose obligations or confer any benefits upon our regulated entities (the inmates) or the public.

Administrative Procedure Act

Because procedures relating to agency management are exempt from the rulemaking provisions of the Administrative Procedure Act (5 U.S.C. 553), we are publishing this change as an interim final rule.

The Administrative Procedure Act (5 U.S.C. 553) allows exceptions to notice-and-comment rulemaking for ''(A) interpretive rules, general statements of policy, or rules of agency organization, procedure, or practice; or (B) when the agency for good cause finds * * * that notice and public procedure thereon are impracticable, unnecessary, or contrary to the public interest.''

This rulemaking is exempt from normal notice-and-comment procedures because these rules are general statements of policy and relate only to internal agency procedure and practice.