Bacitracin methylene disalicylate in grams per ton	Combination in grams per ton	Indications for use	Limitations	Sponsor
* *	*	*	* *	*
(vi) * * *.				
.....	.....	Replacement chickens; as an aid in the prevention of necrotic enteritis caused or complicated by *Clostridium spp. or other organisms susceptible to bacitracin*.	Feed continuously as sole ration..	046573
* *	*	*	* *	*
(ix) * * *.				
.....	.....	Replacement chickens; as an aid in the control of necrotic enteritis caused or complicated by *Clostridium* spp. or other organisms susceptible to bacitracin.	Feed continuously as sole ration. Start at first clinical signs of disease, vary dosage based on severity of infection, administer continuously for 5 to 7 days or as long as clinical signs persist, then reduce medication to prevention level (50 g/t).	046573
* *	*	*	* *	*

* * * * *

Dated: July 10, 1998.

Margaret Ann Miller,

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

[FR Doc. 98–20466 Filed 7–30–98; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 814

[Docket No. 98N–0171]

Medical Devices; Humanitarian Use of Devices

AGENCY: Food and Drug Administration, HHS.

ACTION: Direct final rule; withdrawal.

SUMMARY: The Food and Drug Administration (FDA) published, in the **Federal Register** of April 17, 1998 (63 FR 19185), a direct final rule to implement the amendments to the humanitarian use devices provision of the Federal Food, Drug, and Cosmetic Act (the act), as amended by the Food and Drug Administration Modernization Act of 1997 (FDAMA). The comment period closed July 1, 1998. FDA is withdrawing the direct final rule because the agency received significant adverse comment.

DATES: The direct final rule published at 63 FR 19185, April 17, 1998, is withdrawn effective July 31, 1998.

FOR FURTHER INFORMATION CONTACT: Joanne R. Less, Center for Devices and Radiological Health (HFZ–403), Food and Drug Administration, 9200 Corporate Blvd., Rockville, MD 20857, 301–594–1190.

Therefore, under the Federal Food, Drug, and Cosmetic Act, and under authority delegated to the Commissioner of Food and Drugs, the direct final rule published on April 17, 1998, at 63 FR 19185 is withdrawn.

Dated: July 29, 1998.

William K. Hubbard,

Associate Commissioner for Policy Coordination.

[FR Doc. 98–20594 Filed 7–30–98; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF THE INTERIOR

Office of Surface Mining Reclamation and Enforcement

30 CFR Part 917

[KY–217–FOR]

Kentucky Regulatory Program

AGENCY: Office of Surface Mining Reclamation and Enforcement (OSM), Interior.

ACTION: Final rule; approval of amendment.

SUMMARY: OSM is approving a proposed amendment to the Kentucky regulatory program (hereinafter referred to as the ''Kentucky program'') under the Surface Mining Control and Reclamation Act of 1977 (SMCRA). Kentucky requested the removal of 30 CFR 917.17(a) which disapproved Kentucky's proposed revision to its staffing and budget levels (49 FR 50718, December 31, 1984). The amendment is intended to revise the Kentucky program to be consistent with the Federal regulations and SMCRA.

EFFECTIVE DATE: July 31, 1998.

FOR FURTHER INFORMATION CONTACT: William J, Kovacic, Director, Lexington Field Office, 2675 Regency Road, Lexington, Kentucky 40502. Telephone: (606) 233–2494.

SUPPLEMENTARY INFORMATION:

I. Background on the Kentucky Program

On May 18, 1982, the Secretary of the Interior conditionally approved the Kentucky program. Background information on the Kentucky program, including the Secretary's findings, the disposition of comments, and the conditions of approval can be found in the May 18, 1982 **Federal Register** (47 FR 21404). Subsequent actions concerning conditions of approval and program amendments can be found at