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
* | * | * | * | *

DATES: The direct final rule published at 63 FR 19185, April 17, 1998, is withdrawn effective July 31, 1998.

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.

DEPARTAMENT 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.

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.

SUPPLEMENTARY INFORMATION:
I. Background on the Kentucky Program
II. Submission of the Proposed Amendment
III. Director’s Findings
IV. Summary and Disposition of Comments
V. Director’s Decision
VI. Procedural Determinations

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...