

degree between States due to differences in State laws, the cost of doing business, competition, and other variables.

(iv) NRCS will review payment rates annually, or more frequently as needed, and adjust the rates based upon data from existing contracts, Federal cost rates, and other appropriate sources.

(v) NRCS may adjust payment rates, as needed, on a case-by-case basis, in response to unusual conditions or unforeseen circumstances in delivering technical services such as highly complex technical situations, emergency conditions, serious threats to human health or the environment, or major resource limitations. In these cases, NRCS will set a case-specific not-to-exceed payment rate based on the Department's determination of the scope, magnitude, and timeliness of the technical services needed.

(3) Cost share credits. In order to encourage competitive pricing, a program participant may earn credits toward their cost-share for practice installation under a program contract when a participant selects a technical service provider with prices below the not-to-exceed rates for the provision of technical services. The credits earned will be equal to a percentage of the savings generated by the participant by choosing a lower cost technical service provider. However, in no cases may the application of cost share credits to a program contract result in the Department exceeding any statutory limitations on cost sharing or payments for a particular program.

3. Section 652.1 is amended by revising the definition of *technical service provider* to read as follows:

§ 652.1 Definitions.

* * * * *

Technical service provider means an individual, entity, or public agency certified by NRCS and placed on the approved list to provide technical services to program participants or to the Department.

4. Section 652.4 is amended by adding a new paragraph (h) to read as follows:

§ 652.4 Technical service standards.

* * * * *

(h) Technical service providers may utilize the services of subcontractors to provide specific technical services or expertise needed by the technical service provider, provided that the subcontractors are certified by NRCS in accordance with this part for the particular technical services to be provided and the technical services are provided in terms of their certification agreement. Payments will not be made

for any technical services provided by uncertified subcontractors.

5. In § 652.21 paragraphs (f) and (g) are revised to read as follows:

* * * * *

(f) An individual, private-sector entity, or public agency is conditionally certified provided they had entered into a contract, cooperative agreement, or contribution agreement with the Department prior to March 24, 2003 to provide technical services and they submit an Application for Certification by June 1, 2003. An individual, private-sector entity, or public agency with conditional certification status under this paragraph may continue to provide technical services in accordance with the terms and conditions of the above-described contract, cooperative agreement, or contribution agreement. Conditional certification shall expire either by the date NRCS and the individual, private-sector entity, or public agency enter into a Certification Agreement, as described in § 652.22(c)(1) or September 30, 2003, whichever is earlier.

(g) An individual is conditionally certified if the individual was certified under NRCS policy in effect prior to March 24, 2003, and submits an Application for Certification by June 1, 2003. An individual with conditional certification status under this paragraph may continue to provide technical services to the Department and to program participants in accordance with the above-described prior certification. Conditional certification shall expire either by the date NRCS and the individual enter into a Certification Agreement, as described in § 652.22(c)(1) or September 30, 2003, whichever is earlier.

Signed in Washington, DC on March 7, 2003.

Bruce I. Knight,

Chief, Natural Resources Conservation Service.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 530

[Docket No. 03N-0024]

New Animal Drugs; Phenylbutazone; Extralabel Animal Drug Use; Order of Prohibition; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; correction.

SUMMARY: The Food and Drug Administration (FDA) is correcting a final rule that appeared in the **Federal Register** of February 28, 2003 (68 FR 9528). The document issued an order prohibiting the extralabel use of phenylbutazone animal and human drugs in female dairy cattle 20 months of age or older. FDA is correcting the regulation listing the prohibition by replacing "Phenylbutazone" with "Phenylbutazone in female dairy cattle 20 months of age or older." This correction is being made so that the phenylbutazone listing accurately reflects the agency's intent, which is reflected in the preamble to the final rule.

DATES: This rule is effective May 29, 2003.

FOR FURTHER INFORMATION CONTACT: Gloria J. Dunnava, Center for Veterinary Medicine (HFV-230), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301-827-1168, e-mail: gdunnava@cvm.fda.gov.

SUPPLEMENTARY INFORMATION: In FR Doc. 03-4741, appearing on page 9528 in the **Federal Register** of Friday, February 28, 2003, the following correction is made:

§ 530.41 [Corrected]

On page 9530, in the first column, in § 530.41 *Drugs prohibited for extralabel use in animals*, in paragraph (a)(12), "Phenylbutazone." is corrected to read "Phenylbutazone in female dairy cattle 20 months of age or older."

Dated: March 13, 2003.

Stephen F. Sundlof,

Director, Center for Veterinary Medicine.

[FR Doc. 03-6891 Filed 3-21-03; 8:45 am]

BILLING CODE 4160-01-S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 888

[Docket No. 00N-0018]

Medical Devices; Reclassification of the Knee Joint Patellofemoral Tibial Metal/Polymer Porous-Coated Uncemented Prosthesis and the Knee Joint Femoral Tibial (Unicompartamental) Metal/Polymer Porous-Coated Uncemented Prosthesis

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.