§ 95.6440  Alaska VOR Federal Airway V440 Is Amended to Read in Part

WINOR, AK FIX .................................................................
*FRIDA, AK FIX ............................................................. #10000

§ 95.7001  Jet Routes  § 95.7537  Jet Route J537 Is Amended to Read in Part

ROME, OR VOR/DME ..................................................... MULLAN PASS, ID VOR/DME ........................................ 22000 45000

DEPARTMENT OF HEALTH AND HUMAN SERVICES
Food and Drug Administration

§ 558.500  Ractopamine.

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(e) * * *

(1) * * *

Ractopamine in grams/ton

<table>
<thead>
<tr>
<th>Combination in grams/ton</th>
<th>Indications for use</th>
<th>Limitations</th>
<th>Sponsor</th>
</tr>
</thead>
<tbody>
<tr>
<td>(i) 4.5 to 9 ...............</td>
<td>For increased rate of weight gain, improved feed efficiency, and increased carcass leanness in finishing swine, weighing not less than 150 lbs, led a complete ration containing at least 16% crude protein for the last 45 to 90 lbs of gain prior to slaughter.</td>
<td>Feed continuously as sole ration.</td>
<td>000986</td>
</tr>
</tbody>
</table>

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is classifying norovirus serological reagents into class II (special controls). The special control that will apply to these devices is the guidance document entitled “Class II Special Controls Guidance Document: Norovirus Serological Reagents.” The Agency is classifying these devices into class II (special controls) because special controls, in addition to general controls, will provide a reasonable assurance of safety and effectiveness of these devices and there is sufficient information to establish special controls.

DATES: Effective Date: April 9, 2012. The classification was effective February 23, 2011.

FOR FURTHER INFORMATION CONTACT: Steven Gitterman, Center for Devices and Radiological Health, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 66, Rm. 5518, Silver Spring, MD 20993–0002, 301–796–6694.

SUPPLEMENTARY INFORMATION:
I. Legal Authority

In accordance with section 513(f)(1) of the Federal Food, Drug, and Cosmetic Act (FD&C Act) (21 U.S.C. 360c(f)(1)), devices that were not in commercial distribution before May 28, 1976, the date of enactment of the Medical Device Amendments of 1976 (the 1976...