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 510

Administrative practice and procedure, Animal drugs, Labeling, Reporting and recordkeeping requirements.

- 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 510 is amended as follows:

### PART 510—NEW ANIMAL DRUGS

1. The authority citation for 21 CFR part 510 continues to read as follows:

**Authority:** 21 U.S.C. 321, 331, 351, 352, 353, 360b, 371, 379e.

2. In §510.600, in the table in paragraph (c)(1), remove the entry for “Minrad, Inc.” and alphabetically add a new entry for “Piramal Critical Care, Inc.”; and in the table in paragraph (c)(2), revise the entry for “068507” to read as follows:

#### §510.600 Names, addresses, and drug labeler codes of sponsors of approved applications.

<table>
<thead>
<tr>
<th>Firm name and address</th>
<th>Drug labeler code</th>
</tr>
</thead>
<tbody>
<tr>
<td>Piramal Critical Care, Inc., 3950 Schelden Circle, Bethlehem, PA 18017</td>
<td>068507</td>
</tr>
<tr>
<td>* * * * * * * *</td>
<td></td>
</tr>
</tbody>
</table>

(2) * * *

#### Drug labeler code

<table>
<thead>
<tr>
<th>Firm name and address</th>
<th>Drug labeler code</th>
</tr>
</thead>
<tbody>
<tr>
<td>068507 Piramal Critical Care, Inc., 3950 Schelden Circle, Bethlehem, PA 18017</td>
<td>* * * * * *</td>
</tr>
</tbody>
</table>

Dated: April 15, 2010.

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

### DEPARTMENT OF HEALTH AND HUMAN SERVICES

**Food and Drug Administration**

**21 CFR Part 510**

[Docket No. FDA–2010–N–0002]

**New Animal Drugs; Change of Sponsor’s Name and Address**

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Final rule.

**SUMMARY:** The Food and Drug Administration (FDA) is amending the animal drug regulations to reflect a change of sponsor’s name from Parnell Laboratories (Aust) Pty. Ltd. to Parnell Technologies Pty. Ltd. In addition, the sponsor’s mailing address will be changed.

**DATES:** This rule is effective April 20, 2010.

**FOR FURTHER INFORMATION CONTACT:**

David R. Newkirk, Center for Veterinary Medicine (HFV–100), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 240–276–8307, e-mail: david.newkirk@fda.hhs.gov.

**SUPPLEMENTARY INFORMATION:**

Parnell Laboratories (Aust) Pty. Ltd., Century Estate, unit 6, 476 Gardeners Rd., Alexandria, New South Wales 2015, Australia, has informed FDA that it has changed its name and address to Parnell Technologies Pty. Ltd., unit 4, 476 Gardeners Rd., Alexandria, New South Wales 2015, Australia. Accordingly, the agency is amending the regulations in 21 CFR 510.600(c) to reflect this change.

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 510

Administrative practice and procedure, Animal drugs, Labeling, Reporting and recordkeeping requirements.

- 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 510 is amended as follows:

### PART 510—NEW ANIMAL DRUGS

1. The authority citation for 21 CFR part 510 continues to read as follows:

**Authority:** 21 U.S.C. 321, 331, 351, 352, 353, 360b, 371, 379e.

2. In §510.600, in the table in paragraph (c)(1), revise the entry for “Parnell Laboratories (Aust) Pty. Ltd.”; and in the table in paragraph (c)(2), revise the entry for “068504” to read as follows:

#### §510.600 Names, addresses, and drug labeler codes of sponsors of approved applications.

<table>
<thead>
<tr>
<th>Firm name and address</th>
<th>Drug labeler code</th>
</tr>
</thead>
<tbody>
<tr>
<td>Parnell Technologies Pty. Ltd., unit 4, 476 Gardeners Rd., Alexandria, New South Wales 2015, Australia</td>
<td>068504</td>
</tr>
<tr>
<td>* * * * * * * *</td>
<td></td>
</tr>
</tbody>
</table>

Dated: April 15, 2010.

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

### DEPARTMENT OF HOMELAND SECURITY

**Coast Guard**

**33 CFR Part 165**

[Docket No. USCG–2009–0370]

**RIN 1625–AA11**

**Regulated Navigation Areas; Port of Portland Terminal 4, Willamette River, Portland, OR**

**AGENCY:** Coast Guard, DHS.

**ACTION:** Final rule.

**SUMMARY:** The Coast Guard is establishing two Regulated Navigation Areas (RNA) at the Port of Portland Terminal 4 on the Willamette River in...