in any one year.'' The current threshold after adjustment for inflation is $115 million, using the most current (2003) Implicit Price Deflator for the Gross Domestic Product. FDA does not expect this final rule to result in any 1-year expenditure that would meet or exceed this amount.

VI. Paperwork Reduction Act of 1995

The final rule contains no collections of information. Therefore, clearance by the Office of Management and Budget under the Paperwork Reduction Act of 1995 is not required.

List of Subjects in 21 CFR Part 20

Confidential business information, Courts, Freedom of information, Government employees.

■ Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs, 21 CFR part 20 is amended as follows:

PART 20—PUBLIC INFORMATION

■ 1. The authority citation for part 20 continues to read as follows:

Authority: 5 U.S.C. 552; 18 U.S.C. 1905; 19 U.S.C. 2531–2582; 21 U.S.C. 321–393, 1401–1403; 42 U.S.C. 241, 242, 242a, 242l, 242n, 243, 262, 263, 263b–263n, 264, 265, 300u–300u–5, 300aa–1.

■ 2. Section 20.65 is added to read as follows:

§ 20.65 National defense and foreign policy.

(a) Records or information may be withheld from public disclosure if they are:

(1) Specifically authorized under criteria established by an Executive order to be kept secret in the interest of national defense or foreign policy; and

(2) In fact properly classified under such Executive order.

(b) [Reserved]

■ 3. Section 20.66 is added to read as follows:

§ 20.66 Internal personnel rules and practices.

Records or information may be withheld from public disclosure if they are related solely to the internal personnel rules and practices of the Food and Drug Administration (FDA). Under this exemption, FDA may withhold records or information about routine internal agency practices and procedures. Under this exemption, the agency may also withhold internal records whose release would help some persons circumvent the law.

■ 4. Section 20.67 is added to read as follows:

§ 20.67 Records exempted by other statutes.

Records or information may be withheld from public disclosure if a statute specifically allows the Food and Drug Administration (FDA) to withhold them. FDA may use another statute to justify withholding records and information only if it absolutely prohibits disclosure, sets forth criteria to guide our decision on releasing material, or identifies particular types of matters to be withheld.

■ 5. Section 20.82 is amended by revising paragraph (b)(3) to read as follows:

§ 20.82 Discretionary disclosure by the Commissioner.

* * * * *

(b) * * *

(3) Prohibited from public disclosure under statute.

* * * * *

Dated: July 13, 2005.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. 05–14320 Filed 7–20–05; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 101

Change of Address; Technical Amendment

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; technical amendment.

SUMMARY: The Food and Drug Administration (FDA) is amending its regulations to correct an incorrect address for the Center for Food Safety and Applied Nutrition (CFSAN). This action is editorial in nature and is intended to improve the accuracy of the agency's regulations.

DATES: This rule is effective July 21, 2005.

FOR FURTHER INFORMATION CONTACT: Joyce Strong, Office of Policy and Planning (HF–27), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–7010.

SUPPLEMENTARY INFORMATION: FDA is amending its regulations in § 101.83 (21 CFR 101.83) to reflect the correct address for CFSAN.

Publication of this document constitutes final action on these changes under the Administrative Procedure Act (5 U.S.C. 553). Notice and public procedure are unnecessary because FDA is merely correcting nonsubstantive errors.

List of Subjects in 21 CFR Part 101

Food labeling, Nutrition, Reporting and recordkeeping requirements.

■ Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs, 21 CFR part 101 is amended as follows:

PART 101—FOOD LABELING

■ 1. The authority citation for 21 CFR part 101 continues to read as follows:

Authority: 15 U.S.C. 1453, 1454, 1455; 21 U.S.C. 321, 331, 342, 343, 348, 371; 42 U.S.C. 243, 264, 271.

§ 101.83 [Amended]

■ 2. Section 101.83 is amended in paragraph (c)(2)(ii)(*A*)(*2*) by removing ''200 C St. SW., rm. 2831, Washington, DC 20204'' and by adding in its place ''5100 Paint Branch Pkwy., College Park, MD 20740'' and in paragraph (c)(2)(ii)(*B*)(*2*)by removing ''200 C St., SW., rm. 2831, Washington, DC 20204'' and ''200 C St., SW., Washington DC'' and by adding in their place ''5100 Paint Branch Pkwy., College Park, MD 20740''.

Dated: July 14, 2005.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. 05–14328 Filed 7–20–05; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 558

New Animal Drugs for Use in Animal Feeds; Roxarsone; Semduramycin

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; technical amendment.

SUMMARY: The Food and Drug Administration (FDA) is correcting the single-ingredient roxarsone Type A medicated article that may be used to formulate three-way, combination drug Type C medicated broiler chicken feeds containing semduramicin, virginiamycin, and roxarsone under a new animal drug application (NADA) recently approved for Phibro Animal Health. FDA is also amending the animal drug regulations to reflect two roxarsone Type A medicated articles

approved under separate new animal drug applications (NADAs) for different conditions of use. This action is being taken to improve the accuracy of the agency's regulations.

DATES: This rule is effective July 21, 2005.

FOR FURTHER INFORMATION CONTACT: George K. Haibel, Center for Veterinary Medicine (HFV–6), Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855, 301–827–4567, e-mail: *ghaibel@cvm.fda.gov*.

SUPPLEMENTARY INFORMATION: FDA has found that the list of approved, single-ingredient Type A medicated articles used to formulate three-way, combination drug Type C medicated broiler chicken feeds containing semduramicin, virginiamycin, and roxarsone under NADA 141–226 is in error. The **Federal Register** document that described approval of that application for Phibro Animal Health on February 23, 2004 (69 FR 13221, March 22, 2004), listed 3–NITRO (roxarsone) Type A Medicated article as the source of roxarsone; however, the correct source for this combination feed is ROXARSONE (roxarsone) Type A Medicated article, approved under NADA 92–953. At this time, FDA is amending the regulations in 21 CFR 558.555 to reflect the roxarsone Type A medicated article approved for this combination and a current tabular format.

In addition, FDA has found that the April 1, 2004, edition of parts 500 to 599 (21 CFR parts 500 to 599) of Title 21 of the Code of Federal Regulations (CFR) does not accurately reflect the approved conditions of use for roxarsone Type A medicated articles. Roxarsone is approved as single-ingredient Type A medicated articles under two separate applications, NADA 7–891 for 3–NITRO and NADA 92–953 for ROXARSONE, held by Alpharma, Inc. In error, portions of the regulation describing approvals had been consolidated in July 2000 (65 FR 45711, July 25, 2000). At this time, FDA is amending the regulations in § 558.530 to reflect two separate approvals for roxarsone Type A medicated articles with different approved conditions of use and a current tabular format.

Also, FDA has found that the approved conditions of use codified for NADA 92–953 prior to the July 2000 change were in error. A specific technical amendment to remove turkeys as an approved species (49 FR 30927, August 2, 1984) was reversed in a subsequent change that implemented revised terminology for feed premixes (51 FR 7400, March 3, 1986). At this time, FDA is amending the regulations in § 558.530 to reflect approval of NADA 92–953 for chickens only.

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 558

Animal drugs, Animal feeds.

■ 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 558 is amended as follows:

PART 558—NEW ANIMAL DRUGS FOR USE IN ANIMAL FEEDS

■ 1. The authority citation for 21 CFR part 558 continues to read as follows:

Authority: 21 U.S.C. 360b, 371.

■ 2. Section 558.530 is amended by revising paragraphs (a) through (d)(3); removing paragraph (d)(4); and by redesignating paragraph (d)(5) as paragraph (d)(4) to read as follows:

§ 558.530 Roxarsone.

(a) *Specifications*. Type A medicated articles containing 10, 20, 50, or 80 percent roxarsone.

(b) *Approvals*. See sponsors in § 510.600(c) of this chapter for use as in paragraph (d) of this section.

(1) No. 046573 for use of 10, 20, and 50 percent Type A medicated articles as in paragraph (d)(1)(i) of this section.

(2) No. 046573 for use of 10, 20, 50, and 80 percent Type A medicated articles as in paragraphs (d)(1) through (d)(3) of this section.

(c) *Related tolerances*. See § 556.60 of this chapter.

(d) *Conditions of use*—(1) *Chickens*. It is used in chicken feed as follows:

Roxarsone in grams per ton	Combinations in grams per ton	Indications for use	Limitations	Sponsor
(i) 22.7 to 45.4		Growing chickens: For increased rate of weight gain, improved feed efficiency, and improved pigmentation.	Feed continuously throughout growing period; do not feed to chickens producing eggs for human consumption; withdraw 5 days before slaughter; as sole source of organic arsenic; drug overdose or lack of water may result in leg weakness.	046573
(ii) 22.7 to 45.4	Chlortetracycline 10 to 50	Growing chickens: As in paragraph (d)(1)(i) of this section.	As in paragraph (d)(1)(i) of this section. Chlortetracycline as provided by No. 046573 in § 510.600(c) of this chapter.	
(iii) 22.7 to 45.4	Chlortetracycline 100 to 200	Growing chickens: As in paragraph (d)(1)(i) of this section; and for control of infectious synovitis caused by *Mycoplasma synoviae* susceptible to chlortetracycline.	As in paragraph (d)(1)(i) of this section except feed continuously for 7 to 14 days. Chlortetracycline as provided by No. 046573 in § 510.600(c) of this chapter.	

Roxarsone in grams per ton	Combinations in grams per ton	Indications for use	Limitations	Sponsor
(iv) 22.7 to 45.4	Chlortetracycline 200 to 400	Growing chickens: As in paragraph (d)(1)(i) of this section; and for control of chronic respiratory disease (CRD) and air sac infection caused by *M. gallisepticum* and *Escherichia coli* susceptible to chlortetracycline.	As in paragraph (d)(1)(i) of this section except feed continuously for 7 to 14 days. Chlortetracycline as provided by No. 046573 in § 510.600(c) of this chapter.	
(v) 22.7 to 45.4	Chlortetracycline 500	Growing chickens: As in paragraph (d)(1)(i) of this section; and for reduction of mortality due to *E. coli* infections susceptible to chlortetracycline.	As in paragraph (d)(1)(i) of this section except feed continuously for 5 days. Chlortetracycline as provided by No. 046573 in § 510.600(c) of this chapter.	

(2) *Turkeys.* It is used in turkey feed as follows:

Roxarsone in grams per ton	Combinations in grams per ton	Indications for use	Limitations	Sponsor
(i) 22.7 to 45.4		Growing turkeys: For increased rate of weight gain, improved feed efficiency, and improved pigmentation.	Feed continuously throughout growing period; do not feed to turkeys producing eggs for human consumption; withdraw 5 days before slaughter; as sole source of organic arsenic; drug overdose or lack of water may result in leg weakness.	046573
(ii) 22.7 to 45.4	Chlortetracycline 10 to 50	Growing turkeys: As in paragraph (d)(2)(i) of this section.	As in paragraph (d)(2)(i) of this section. Chlortetracycline as provided by No. 046573 in § 510.600(c) of this chapter.	
(iii) 22.7 to 45.4	Chlortetracycline 200	Growing turkeys: As in paragraph (d)(2)(i) of this section; and for control of infectious synovitis caused by *Mycoplasma synoviae* susceptible to chlortetracycline.	As in paragraph (d)(2)(i) of this section except feed continuously for 7 to 14 days. Chlortetracycline as provided by No. 046573 in § 510.600(c) of this chapter.	
(iv) 22.7 to 45.4	Chlortetracycline 400	1. Growing turkeys: As in paragraph (d)(2)(i) of this section; and for control of hexamitiasis caused by *Hexamita meleagrides* susceptible to chlortetracycline. 2. Turkey poults not over 4 weeks of age: Reduction of mortality due to paratyphoid caused by *Salmonella typhimurium* susceptible to chlortetracycline.	As in paragraph (d)(2)(i) of this section except feed continuously for 7 to 14 days. Chlortetracycline as provided by No. 046573 in § 510.600(c) of this chapter.	
(v) 22.7 to 45.4	Chlortetracycline 25 mg/lb body weight daily	Growing turkeys: As in paragraph (d)(2)(i) of this section; and for control of complicating bacterial organisms associated with bluecomb (transmissible enteritis, coronaviral enteritis) susceptible to chlortetracycline.	As in paragraph (d)(2)(i) of this section except feed continuously for 7 to 14 days. Chlortetracycline as provided by No. 046573 in § 510.600(c) of this chapter.	

(3) *Swine.* It is used in swine feed as follows:

Roxarsone in grams per ton	Combinations in grams per ton	Indications for use	Limitations	Sponsor
(i) 22.7 to 34.1		Growing and finishing swine: For increased rate of weight gain and improved feed efficiency.	Feed continuously throughout growing period; withdraw 5 days before slaughter; as sole source of organic arsenic.	046573
(ii) 22.7 to 34.1	Chlortetracycline 400 (to administer 10 mg/lb body weight)	Growing and finishing swine: As in paragraph (d)(3)(i) of this section; and for treatment of bacterial enteritis caused by *E. coli* and *S. choleraesuis* and bacterial pneumonia caused by *Pasteurella multocida* susceptible to chlortetracycline.	Feed for not more than 14 days; withdraw 5 days before slaughter; as sole source of organic arsenic.	
(iii) 181.5		Growing and finishing swine: For the treatment of swine dysentery.	Feed for not more than 6 consecutive days; if improvement is not observed, consult a veterinarian; withdraw 5 days before slaughter; as a sole source of organic arsenic; animals must consume enough medicated feed to provide a therapeutic dose.	046573
(iv) 181.5	Chlortetracycline 10 to 50	Growing and finishing swine: As in paragraph (d)(3)(i) of this section; and for treatment of swine dysentery.	Feed for not more than 6 consecutive days; if improvement is not observed, consult a veterinarian; withdraw 5 days before slaughter; as a sole source of organic arsenic; animals must consume enough medicated feed to provide a therapeutic dose.	
(v) 181.5	Chlortetracycline 400 (to administer 10 mg/lb body weight)	Growing and finishing swine: As in paragraph (d)(3)(iii) of this section; and for treatment of bacterial enteritis caused by *E. coli* and *S. choleraesuis* and bacterial pneumonia caused by *P. multocida* susceptible to chlortetracycline.	Feed for not more than 6 consecutive days; if improvement is not observed, consult a veterinarian; withdraw 5 days before slaughter; as a sole source of organic arsenic; animals must consume enough medicated feed to provide a therapeutic dose.	

* * * * *

■ 3. Section 558.555 is amended by revising paragraph (d) to read as follows:

§ 558.555 Semduramicin.

* * * * *

(d) *Conditions of use in chickens.* It is used in chicken feed as follows:

Semduramicin in grams per ton	Combinations in grams per ton	Indications for use	Limitations	Sponsor
(1) 22.7 (25 ppm)		Broiler chickens: For the prevention of coccidiosis caused by *Eimeria acervulina*, *E. brunetti*, *E. maxima*, *E. mivati/E. mitis*, *E. necatrix*, and *E. tenella*.	Do not feed to laying hens.	066104
(2) 22.7	Bacitracin methylene disalicylate 10 to 50	Broiler chickens: As in paragraph (d)(1) of this section; for improved feed efficiency.	Feed continuously as sole ration. Do not feed to laying hens. Bacitracin methylene disalicylate as provided by No. 046573 in § 510.600(c) of this chapter.	066104

Semduramicin in grams per ton	Combinations in grams per ton	Indications for use	Limitations	Sponsor
(3) 22.7	Bacitracin methylene disalicylate 10 to 50 plus roxarsone 45.4	Broiler chickens: As in paragraph (d)(4) of this section; for improved feed efficiency.	Feed continuously as sole ration. Use feed within 2 weeks of production. Do not feed to laying hens. Use as sole source of organic arsenic. Poultry should have access to drinking water at all times. Drug overdosage or lack of water intake may result in leg weakness or paralysis. Withdraw 5 days before slaughter. Bacitracin methylene disalicylate and roxarsone as provided by No. 046573 in § 510.600(c) of this chapter.	066104
(4) 22.7	Roxarsone 45.4	Broiler chickens: For the prevention of coccidiosis caused by *Eimeria acervulina*, *E. brunetti*, *E. maxima*, *E. mivati/E. mitis*, *E. necatrix*, and *E. tenella*, including some field strains of *E. tenella* that are more susceptible to semduramicin combined with roxarsone than semduramicin alone.	Feed continuously as sole ration. For broiler chickens only. Do not feed to laying hens. Use as sole source of organic arsenic. Withdraw 5 days before slaughter. Roxarsone as provided by No. 046573 in § 510.600(c) of this chapter.	066104
(5) 22.7	Virginiamycin 5	Broiler chickens: As in paragraph (d)(1) of this section; for increased rate of weight gain and improved feed efficiency.	Feed continuously as sole ration. Do not feed to laying hens. Virginiamycin as provided by No. 066104 in § 510.600(c) of this chapter.	066104
(6) 22.7	Virginiamycin 5 to 15	Broiler chickens: As in paragraph (d)(1) of this section; for increased rate of weight gain.	Feed continuously as sole ration. Do not feed to laying hens. Virginiamycin as provided by No. 066104 in § 510.600(c) of this chapter.	066104
(7) 22.7	Virginiamycin 20	Broiler chickens: As in paragraph (d)(1) of this section; for prevention of necrotic enteritis caused by *Clostridium perfringens* susceptible to virginiamycin.	Feed continuously as sole ration. Do not feed to laying hens. Virginiamycin as provided by No. 066104 in § 510.600(c) of this chapter.	066104
(8) 22.7	Virginiamycin 20 plus roxarsone 22.7 to 45.4	Broiler chickens: As in paragraph (d)(1) of this section; for prevention of necrotic enteritis caused by *Clostridium perfringens* susceptible to virginiamycin; for increased rate of weight gain and improved feed efficiency; and for improved pigmentation.	Feed continuously as sole ration throughout growing period. Do not feed to laying hens. Use as sole source of organic arsenic. Poultry should have access to drinking water at all times. Drug overdose or lack of water may result in leg weakness. Roxarsone as in § 558.530(b)(1) of this chapter provided by No. 046573 in § 510.600(c) of this chapter; semduramicin and virginiamycin as provided by No. 066104.	066104

Dated: April 25, 2005.

Steven D. Vaughn,

Director, Office of New Animal Drug Evaluation.

[FR Doc. 05–14329 Filed 7–20–05; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF THE INTERIOR

Bureau of Indian Affairs

25 CFR Part 124

RIN 1076–AE74

Deposit of Proceeds From Lands Withdrawn for Native Selection; Correction

AGENCY: Bureau of Indian Affairs, Interior.

ACTION: Final rule; correction.

SUMMARY: This document contains a correction to a final rule that was published Thursday, July 14, 2005 (70 FR 40660). The regulation relates to Deposit of Proceeds from Lands Withdrawn for Native Selection.

EFFECTIVE DATE: July 14, 2005.

FOR FURTHER INFORMATION CONTACT: Assistant Director, Office of Trust Regulations, Policies and Procedures, by telephone at (505) 816–1086, or by facsimile transmission at (505) 816–1377.

SUPPLEMENTARY INFORMATION: This rule is published by the authority of the Secretary, granted under 43 U.S.C. 1601 *et seq.* and 25 U.S.C. 4001 *et seq.*, and delegated to the Assistant Secretary—Indian Affairs 209 DM 8.1.

Background

The final rule provides contact information to be used by all Departments and Agencies, the State of Alaska, and any other interested parties for deposit of proceeds from lands withdrawn for native selection. This rule was published by the Assistant Secretary—Indian Affairs in consultation with the Special Trustee for American Indians under the provisions of the American Indian Trust Fund Management Reform Act of 1994.

Need for Correction

As published, the final rule was introduced by words of issuance that do not satisfy Office of the Federal Register standards. The language must be corrected to allow for correct codification of the revised regulation.

Correction of Publication

Accordingly, the publication on July 14, 2005, of the final rule that was the subject of FR Doc. 05–13891, is corrected as follows:

On page 40660, in the second column, immediately following the name and title of the document's signer, in the words of issuance, the word ''amended'' is corrected read ''revised.''

Dated: July 15, 2005.

James E. Cason,

Associate Deputy Secretary of the Interior.

[FR Doc. 05–14437 Filed 7–20–05; 8:45 am]

BILLING CODE 4310–2W–P

ENVIRONMENTAL PROTECTION AGENCY

40 CFR Part 52

[Docket # ID–03–003; FRL–7941–7]

Approval and Promulgation of Air Quality Implementation Plan; Idaho; Correction

AGENCY: Environmental Protection Agency (EPA).

ACTION: Final rule; correction.

SUMMARY: This document corrects the preamble to a final rule published in the **Federal Register** of July 11, 2005 (70 FR 39658) regarding revisions to the open burning regulations in Idaho's State Implementation Plan. This notice clarifies that, under section 307(b)(1) of the Clean Air Act, any petition for judicial review of this action must be filed in the United States Court of Appeals for the appropriate circuit within 60 days from the date notice of approval appeared in the **Federal Register**, and not 30 days, as erroneously stated in July 11, 2005 action.

FOR FURTHER INFORMATION CONTACT: Donna Deneen, (206) 553–6706.

SUPPLEMENTARY INFORMATION:

Correction

In the final rule, beginning on page 39658 in the issue of July 11, 2005, make the following correction, in the **SUPPLEMENTARY INFORMATION** section. On page 39661 in the 3rd column, remove ''August 10, 2005'' in the first paragraph and replace it with ''September 9, 2005''.

Dated: July 14, 2005.

Michelle Pirzadeh,

Acting Regional Administrator, Region 10.

[FR Doc. 05–14399 Filed 7–20–05; 8:45 am]

BILLING CODE 6560–50–M

ENVIRONMENTAL PROTECTION AGENCY

40 CFR Part 52

[R06–OAR–2005–NM–0001; FRL–7942–5]

Approval and Promulgation of Air Quality Implementation Plans; New Mexico; Albuquerque/Bernalillo County

AGENCY: Environmental Protection Agency (EPA).

ACTION: Final rule.

SUMMARY: This action finalizes our approval of the State Implementation Plan (SIP) revisions submitted by the Governor of New Mexico on September 7, 2004. The submittal revises the second ten-year carbon monoxide (CO) maintenance plan for the Albuquerque/Bernalillo County, New Mexico area. The submittal also revises the relevant parts of the New Mexico Administrative Code (NMAC) including revisions to the General Provisions, Inspection and Maintenance (I&M) Program, and the contingency measures. We are finalizing approval of these revisions in accordance with the requirements of the Federal Clean Air Act (the Act).

DATES: This rule is effective on August 22, 2005.

ADDRESSES: The EPA has established a docket for this action under Regional Material in EDocket (RME) Docket ID No. R06–OAR–2005–NM–0001. All documents in the docket are listed in the RME index at *http://docket.epa.gov/rmepub/,* once in the system, select ''quick search,'' then key in the appropriate RME Docket identification number. Although listed in the index, some information is not publicly available, *i.e.*, CBI or other information whose disclosure is restricted by statute. Certain other material, such as copyrighted material, is not placed on the Internet and will be publicly available only in hard copy form. Publicly available docket materials are available either electronically in RME or in hard copy at the Air Planning Section (6PD–L), Environmental Protection Agency, 1445 Ross Avenue, Suite 700, Dallas, Texas 75202–2733. The file will be made available by appointment for public inspection in the Region 6 FOIA Review Room between the hours of 8:30 a.m. and 4:30 p.m. weekdays except for legal holidays. Contact the person listed in the **FOR FURTHER INFORMATION CONTACT** paragraph below to make an appointment. If possible, please make the appointment at least two working days in advance of your visit. There will be a 15 cent per page fee for making