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(iv) Make arrests without warrant for an offense under the act with respect to counterfeit drugs if the offense is committed in the presence of the criminal investigator or, in the case of a felony, if the investigator has probable cause to believe that the person so arrested has committed, or is committing, such offense; and

(v) Make, prior to the institution of libel proceedings under section 304(a)(2) of the act (21 U.S.C. 334(a)(2)), seizures of drugs or containers or of equipment, punches, dies, plates, stones, labeling, or other things, if they are, or the criminal investigator has reasonable grounds to believe that they are, subject to seizure and condemnation under section 304(a)(2) of the act.

(2) Perform such other functions under the act, or any other law, as the Commissioner may prescribe.

(3) To administer oaths and affirmations under section 1 of the act of January 31, 1925 (Ch. 124, 43 Stat. 803); sections 12 to 15 of Reorganization Plan No. IV, effective June 30, 1940; and Reorganization Plan No. 1 of 1953, effective April 11, 1953.

(c) Any officer or employee of the Food and Drug Administration who has been designated by the Commissioner to provide specialized law enforcement support involving criminal investigations under the act, and other duties as assigned by the Commissioner, and issued the Food and Drug Administration Official Credential consisting of Form FDA-200E, Special Authority for Criminal Investigative Specialists, is authorized to receive information as to all matters relating to such act and regulations issued under the act.

(d) These officials may not further redelegate these authorities.

§5.32 Certification following inspections.

Regional Food and Drug Directors and District Directors are authorized to issue certificates of sanitation under §1240.20 of this chapter. These officials may not further redelegate this authority.

§5.33 Issuance of reports of minor violations.

(a) The following officials are authorized to perform all the functions of the

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Commissioner of Food and Drugs (Commissioner) under section 309 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 336) (the act) regarding the issuance of written notices or warnings:

(1)(i) The Director and Deputy Directors, Center for Biologics Evaluation and Research (CBER).

(ii) The Director and Deputy Directors, Office of Compliance and Biologics Quality, CBER.

(2)(i) The Director and Deputy Directors for Science and for Regulations and Policy, Center for Devices and Radiological Health (CDRH).

(ii) The Director and Deputy Director, Office of Compliance, CDRH.

(iii) For medical devices assigned to their respective divisions, the Division Directors, Office of Compliance, CDRH.

(iv) The Director and Deputy Director, Office of Surveillance and Biometrics (OSB), CDRH, and the Director and Deputy Director, Division of Surveillance Systems (DSS), OSB, CDRH.

(3)(i) The Director and Deputy Director, Center for Food Safety and Applied Nutrition (CFSAN).

(ii) The Director of Regulations and Policy, CFSAN.

(iii) The Director, Office of Field Programs, CFSAN.

(iv) The Director, Division of Enforcement and Programs, Office of Field Programs, CFSAN.

(4)(i) The Director and Deputy Director, Center for Veterinary Medicine (CVM).

(ii) The Director and Deputy Director, Office of Surveillance and Compliance, CVM.

(iii) The Director, Division of Compliance, Office of Surveillance and Compliance, CVM.

(5)(i) The Director, the Deputy Director, the Associate Director for Regulatory Policy, and the Directors, Office of Review Management and Office of Pharmaceutical Science, Center for Drug Evaluation and Research (CDER).

(ii) The Director and Deputy Director, Office of Compliance, CDER.

(iii) The Associate Director for Medical Policy, CDER.

(iv) The Director, Division of Drug Marketing, Advertising, and Communications, Office of Medical Policy, CDER.

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(6)(i) Regional Food and Drug Directors.

(ii) District Directors.

(iii) Chiefs of District Compliance Branches.

(iv) The Director, St. Louis Branch.

(v) The Director, Northeast Regional Laboratory, Northeast Region.

(vi) The Director, Southeast Regional Laboratory, Southeast Region.

(vii) The Director, Winchester Engineering and Analytical Center.

(viii) The Director, National Forensic Chemistry Center.

(ix) The Director, Arkansas Regional Laboratory.

(b) The following officials are authorized to perform all the functions of the Commissioner under section 539(d) of the act (21 U.S.C. 360pp(d)) regarding the issuance of written notices or warnings:

(1) The Director and Deputy Directors for Science and for Regulations and Policy, Center for Devices and Radiological Health (CDRH).

(2) The Director and Deputy Director, Office of Compliance, CDRH.

(3) For medical devices assigned to their respective divisions, the Division Directors, Office of Compliance, CDRH.

(4) The Director and Deputy Director, Office of Surveillance and Biometrics (OSB), CDRH, and the Director and Deputy Director, Division of Surveillance Systems (DSS), OSB, CDRH.

(5) Regional Food and Drug Directors; District Directors; the Director, St. Louis Branch; the Director, Northeast Regional Laboratory, Northeast Region; the Director, Southeast Regional Laboratory, Southeast Region; the Director, Winchester Engineering and Analytical Center; the Director, National Forensic Chemistry Center, and the Director, Arkansas Regional Laboratory when such functions relate to:

(i) Assemblers of diagnostic x-ray systems, as defined in § 1020.30(b) of this chapter; and

(ii) Manufacturers of sunlamp products and ultraviolet lamps intended for use in any sunlamp product as defined in § 1040.20(b) of this chapter.

(c) These officials may not further redelegate these authorities.

§ 5.34 Issuance of notices relating to proposals and orders for debarment and denial of an application to terminate debarment.

(a) The Director, the Deputy Director, and the Associate Director for Regulatory Policy, Center for Drug Evaluation and Research, the Director and Deputy Director, Center for Veterinary Medicine, and the Director and Deputy Directors, Center for Biologics Evaluation and Research are authorized to issue the following notices and make all findings required in relation to these notices under section 306 of the Federal Food, Drug, and Cosmetic Act (the act) (21 U.S.C. 335a) which relate to the assigned functions of their organizations:

(1) Notices of opportunity for hearing on proposals for mandatory or permissive debarment.

(2) Notices ordering debarment when opportunity for a hearing has been waived.

(3) Notices ordering debarment where the person notifies the agency that the person consents to debarment under section 306(c)(2)(B) of the act (21 U.S.C. 335a(c)(2)(B)).

(4) Notices of opportunity for hearing on proposals denying an application to terminate debarment under section 306(d)(3) of the act (21 U.S.C. 335u(d)(3)).

(5) Orders denying an application to terminate debarment under section 306(d)(3) of the act (21 U.S.C. 335u(d)(3)) when opportunity for a hearing has been waived.

(b) These officials may not further redelegate these authorities.

§ 5.35 Officials authorized to make certification under 5 U.S.C. 605(b) for any proposed and final rules.

(a) The following officials are authorized to perform all the functions of the Commissioner of Food and Drugs with regard to decisions made under the Regulatory Flexibility Act (5 U.S.C. 605(b)), to certify that a proposed or final rule, if issued, will not have a significant economic impact on a substantial number of small entities:

(1) The Associate Commissioner for Regulatory Affairs.

(2) The Director, Center for Biologics Evaluation and Research.