

FDA STAFF MANUAL GUIDES, VOLUME II - DELEGATIONS OF AUTHORITY

REGULATORY - ANIMAL DRUGS

ISSUANCE OF FEDERAL REGISTER DOCUMENTS PERTAINING TO THE DETERMINATION OF SAFE LEVELS, NOTICE OF NEED FOR DEVELOPMENT OF AN ANALYTICAL METHOD, NOTICE OF AVAILABILITY OF A DEVELOPED ANALYTICAL METHOD, AND PROHIBITION OF CERTAIN EXTRALABEL DRUG USE

Effective Date: October 16, 2014

1. AUTHORITIES DELEGATED AND TO WHOM DELEGATED.

A. Director and Deputy Directors, Office of Foods and Veterinary Medicine, are authorized to issue Federal Register documents pertaining to the determination of safe levels, notice of need for development of an analytical method, notice of availability of a developed analytical method, and prohibition of certain extralabel drug use related to implementation of sections 512(a)(4) and (5) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360b(a)(4) and (5)).

2. REDELEGATION.

These officials may further re-delegate this authority.

3. EFFECTIVE DATE.

The Commissioner of Food and Drugs approved this delegation on October 16, 2014.

STATUS (I, R, C)	DATE APPROVED	LOCATION OF CHANGE HISTORY	CONTACT	APPROVING OFFICIAL
Initial	02/24/2011	N/a	CVM/OM	Commissioner of Food and Drugs
Revision	10/16/2014	N/a	CVM/OM	Margaret A. Hamburg, M.D. Commissioner of Food and Drugs