

FDA Staff Manual Guides, Volume I – Organizations and Functions

Department of Health and Human Services

Food and Drug Administration

Center for Drug Evaluation and Research

Office of New Drugs

Office Regulatory Operations

Division of Regulatory Operations for Nonprescription Drugs

Effective Date: September 25, 2019

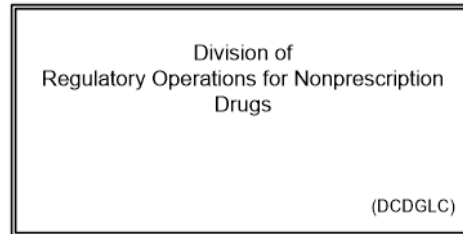
1. Division of Regulatory Operations for Nonprescription Drugs (DCDGLC).

- A. Serves as the regulatory leader on product-specific review teams, providing day-to-day regulatory guidance to review staff on product-specific issues.
- B. Provides office and division leadership with regulatory expertise on non-prescription products and cross-cutting programs, ensuring consistency with legal and regulatory requirements, as well as office-wide policies.
- C. Manages and coordinates regulatory operations for a portfolio of regulated products, from the investigational stage through marketing application review and post-marketing oversight.
- D. Functions as the communication liaison between other offices within Office of New Drugs, centers, Food and Drug Administrations and the regulated industry for product-specific interactions.
- E. Collaborates in the design and implementation of processes and policies as needed to ensure compliance with relevant laws, regulations, and user fee program commitments.

2. Authority and Effective Date.

The functional statements for the Division of Regulatory Operations for Nonprescription Drugs were approved by the Secretary of Health and Human Services on September 25, 2019.

**Department of Health and Human Services
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The following is the Department of Health and Human Services, Food and Drug Administration, Center for Drug Evaluation and Research, Office of New Drugs, Office of Regulatory Operations, Division of Regulatory Operations for Nonprescription Drugs organization structure depicting all the organizational structures reporting to the Director.

Division of Regulatory Operations for Nonprescription Drugs (DCDGLC).