

**FDA Staff Manual Guides, Volume I – Organizations and Functions**

**Department of Health and Human Services**

**Food and Drug Administration**

**Center for Devices and Radiological Health**

**Office of Product Evaluation and Quality**

**Office of Health Technology VI**

**Division of Health Technology VI B**

Effective Date: December 14, 2018

**1. Division of Health Technology VI B (DCCFHB).**

- A. Serves as the primary source for scientific and medical expertise on medical devices with regard to safety and effectiveness.
- B. Carries out scientific and end-to-end medical device review evaluation.
- C. Coordinates actions on classification of medical devices.
- D. Coordinates, carries out, and makes premarket review determinations.
- E. Plans and coordinates post market compliance and enforcement efforts related to medical devices. Participates in development and interpretation of post market regulations and policies related to medical devices.
- F. Reviews and evaluates design, test, and production data and reports from manufacturers to ensure compliance with promulgated standards and regulations.

**2. Authority and Effective Date.**

The functional statements for the Division of Health Technology VI B were approved by the Secretary of Health and Human Services and effective on December 14, 2018.

**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 Devices and Radiological Health, Office of Product Evaluation and Quality, Office of Health Technology VI, Division of Health Technology VI B organization structure depicting all the organizational structures reporting to the Director.

Division of Health Technology VI B (DCCFHB)