

us fda drug products pdf

Original New Drug Approvals (NDAs and BLAs) by Month All applications approved for the first time during the selected month. Includes New Molecular Entities (NMEs) and new biologics.

Drugs@FDA: FDA Approved Drug Products

Drug Shortages can occur for many reasons including manufacturing and quality problems, delays, and discontinuations. Manufacturers provide FDA most drug shortage information, and the agency works ...

Drug Shortages - Food and Drug Administration

Laser Products - Conformance with IEC 60825-1 Ed. 3 and IEC 60601-2-22 Ed. 3.1 (Laser Notice No. 56) - Draft Guidance for Industry and Food and Drug Administration Staff (PDF - 118KB); The FDA ...

Radiation-Emitting Products - Food and Drug Administration

The Food and Drug Administration (FDA or USFDA) is a federal agency of the United States Department of Health and Human Services, one of the United States federal executive departments. The FDA is responsible for protecting and promoting public health through the control and supervision of food safety, tobacco products, dietary supplements, prescription and over-the-counter pharmaceutical drugs ...

Food and Drug Administration - Wikipedia

Points of Contact for Questions Regarding Registration and Listing for Human and Animal Drugs and Biologics; Guidance for Industry: Providing Regulatory Submissions in Electronic Format -Drug Establishment Registration and Drug Listing (PDF - 776 KB)

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