CDER Product-Specific Guidances Withdrawn Listing

Updated April 10th, 2023

Active Ingredient	Type of Guidance	Route and Dosage Form	RLD	Date PSG Posted or Revised	Federal Register Notice / Withdrawal Date
BUTENAFINE HYDROCHLORIDE	Draft	Topical Cream	021408	03/01/2012	02/01/2015
HYDROXYPROGESTERONE CAPROATE	Draft	Subcutaneous Solution	021945	09/16/2019	04/06/2023
LEVONORGESTREL	Draft	Intrauterine Device	203159	04/01/2014	10/01/2014
LORCASERIN HYDROCHLORIDE	Draft	Oral Tablet, Extended Release	022529	03/01/2015	03/04/2021
LORCASERIN HYDROCHLORIDE	Draft	Oral Tablet, Extended Release	208524	05/01/2017	03/04/2021
LOVASTATIN; NIACIN	Draft	Oral Tablet, Extended Release	021249	07/01/2009	04/18/2016
NIACIN; SIMVASTATIN	Draft	Oral Tablet, Extended Release	022078	10/01/2011	04/18/2016