



**OFFICE OF SUPPLY CHAIN MANAGEMENT
Sheriff's Prescription Drug Take Back Program**

Repackager/Retailer Notification Form

Pharmaceutical Repackagers/Retailers¹ subject to the Cook County Safe Disposal Ordinance are required to provide notification to the Director of the Cook County Sheriff's Prescription Drug Take Back Program as to contact information of the Manufacturer from whom the retailer or Repackager obtains a Covered Drug², including the telephone number, mailing address and email address of the retailer's or Repackager's point of contact at the Manufacturer. Said notification to the Director is required on or after April 23, 2017, or by six months after a retailer whose label appears on a Covered Drug or the Covered Drug's packaging starts selling the Covered Drug in the County, or by six months after a Covered Drug repackaged by Repackager is first sold in the County, and, thereafter. Sec. 46-104 (Collection Plan-Participation)

A Repackager has fulfilled its notification requirement when this completed fillable form is sent via email to our email address at sheriff.recycling@cookcountyil.gov.

Please print when completing this form.

Name of Person Completing Form for Repackager/Retailer		Date Form Submitted
Title of Person Completing Form for Repackager/Retailer		Email Address of Person Completing Form
Name of Repackager/Retailer		Repackager's Mailing Address
Name of Covered Drug	Type of Drug	Name of Manufacturer
Point of Contact of Repackager at the Manufacturer		Manufacturer's Telephone Number ()
Manufacturer's Mailing Address, City, State and Zip Code		Manufacturer's Email Address

¹Repackager is defined as a person who owns or operates an establishment that repacks and relabels a product or package for further sale, or for distribution without a further transaction. Sec.46-102

² A Covered Drug means a Drug sold, offered for sale or dispensed in Cook County in any form, including prescription, nonprescription, brand name and generic drugs. Covered Drug does not include (1) vitamins or supplements, (2) herbal-based remedies and homeopathic drugs, products or remedies; 3) cosmetics, shampoos, sunscreens, toothpaste, lip balm, antiperspirants, or other personal care products that are regulated as both cosmetics and nonprescription drugs under the federal Food, Drug and Cosmetic Act (Title 21 U.S.C. Chapter 9); 4) Drugs for which Producers provide a pharmaceutical stewardship or take-back program as part of a federal Food and Drug Administration-managed risk evaluation and mitigation strategy (21 U.S.C. Sec. 355-1); 5) Drugs that are biological products as defined by 21 C.F.R. Sec. 600.3(h) as its exists on the effective date of this Ordinance if the Producer already provides a pharmaceutical product stewardship or take back program; and 6) medical devices or their components parts or accessories. Sec. 46-102.