Vanish (2017-09-22)

Report a Concern

Starting date:	September 22, 2017
Type of communication:	Drug Recall
Subcategory:	Drugs
Hazard classification:	Type III
Source of recall:	Health Canada
Issue:	Product Safety
Audience:	Healthcare Professionals, General Public, Hospitals
Identification number:	RA-64684

- Reason
- Depth of distribution
- Affected products

Affected products

Vanish

Reason

Affected lot was released to market without adequate testing.

Depth of distribution

Wholesalers and end-users (dental offices) in Ontario

Affected products

Vanish

DIN, NPN, DIN-HIM

NPN 80003435

Dosage form

Liquid

Strength

Sodium Fluoride 5.0%

Lot or serial number

N871667

Companies

Recalling Firm	3M Canada Company 300 Tartan Drive London N5V 4M9 Ontario CANADA
Marketing Authorization Holder	3M Canada Company 300 Tartan Drive London N5V 4M9 Ontario CANADA

Date modified: 2017-10-03