Government of Canada Gouvernement du Canada

Canada

Recalls and safety alerts

Viibryd Starter Kit (10 & 20 mg)

Report a Concern

Starting date:	May 7, 2018
Type of communication:	Drug Recall
Subcategory:	Drugs
Hazard classification:	Type II
Source of recall:	Health Canada
Issue:	Product Safety
Audience:	Healthcare Professionals
Identification number:	RA-66756

- Reason
- Depth of distribution
- Affected products

Recalled Products

Viibryd Starter Kit (10 & 20 mg)

Reason

Affected lots are labelled with the incorrect strengths on the blister card.

Depth of distribution

Distributed to doctors in BC, AB, SK, ON, QC, NB, NS

Affected products

Viibryd Starter Kit (10 & 20 mg)

DIN, NPN, DIN-HIM

DIN 02443759

Dosage form

Kit, Tablet

Strength

10 & 20 mg

Lot or serial number

- 10 mg: 1540126, 1540127, 1540128
- 20 mg: 1540126, 1540127, 1540128

Companies

Recalling Firm	Allergan Inc. 85 Enterprise Blvd, Suite 500 Markham L6G 0B5 Ontario CANADA
Marketing Authorization Holder	Forest Laboratories Canada Inc. 500-85 Enterprise Blvd Markham L6G 0B5 Ontario CANADA

Date modified: 2018-05-09