Health Canada Santé Canada

Mandatory Medical Device Problem Reporting Form for Industry

CANADA VIGILANCE - MEDICAL DEVICE PROBLEM REPORTING PROGRAM

If more space is required, please attach additional sheets
Fields required to be completed for updates/final reports are indicated by an *

A. REPORTER INFORMATION

1. i. Reporter Type

(●) Manufacturer ( ) Importer

In the case where the reporter is the importer:

ii. Did the importer report the incident to the manufacturer?

( ) Yes ( ) No

iii. Is the importer also submitting the report on behalf of the manufacturer?

( ) Yes ( ) No

2. Reporter Contact Information *

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3. Reporter File No. *

INF-US-2020-016493 (368385)

4. Health Canada File No. (if applicable) *

5. Type of Report *

( ) Preliminary ( ) Update (●) Final ( ) Preliminary & Final

If "preliminary" only, anticipated date for the final report:

(YYYY-MM-DD)

If "update/final", date the previous report was submitted to Health Canada:

2020-08-14 (YYYY-MM-DD)

6. Date Submitted *

2020-08-14 (YYYY-MM-DD)

	Manufacturer	Importer
7. Name and Address	CAREFUSION 303, INC. 10020 Pacific Mesa Blvd. San Diego, CA 92121-2733	Becton Dickinson Canada Inc. 2100 Derry Road, Suite 100, Mississauga, ON L5N 0B3 Canada
8. Health Canada assigned company identification number (if known):	105302	101291
9. Establishment License Number (if applicable):	N/A	204

B. INCIDENT INFORMATION

1. Classification of Incident *

i. ( ) 10-Day (●) 30-Day

ii. (●) Canadian ( ) Foreign

iii. [ ] Investigational testing [ ] Special Access Program
[ ] Radiation emitting device (if applicable)

2. Date of Incident

(YYYY-MM-DD)

3. Reporter's Awareness Date

(YYYY-MM-DD)

4. Patient Consequences

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5. Details of Incident

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C. MEDICAL DEVICE INFORMATION

1. **Trade/Brand Name ***

2. **Control/Lot/Serial No.**

3. **Expiration Date**

(YYYY-MM-DD)

4. **i. Device Classification**

○ I ○ II ◉ III ○ IV

ii. Device License No.

iii. Device Identification No

iv. Manufacturer's Medical Device Identifier (catalogue/model no.)

5. **Software Version**

6. **Age of Device**

7. **How long was the device in use?**

8. **Was the device labelled as sterile?**

○ Yes ◉ No

9. **Availability of device for evaluation**

○ Destroyed ◉ Returned to Manufacturer/Importer

○ Neither (with explanation)

D. COMPLAINANT INFORMATION

1. **Complainant is a:**

○ Consumer ○ Health professional ◉ Other

2. **Name of Complainant**

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3. **Name of Health Care Facility (if applicable)**

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4. **Address**

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5. **Telephone No. and/or E-mail Address**

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Privacy Notice Statement: For the purposes of the Canada Vigilance - Medical Device Problem Reporting Program, information related to the identity of the complainant and/or reporter will be protected as personal information under the *Privacy Act*, and under the *Access to Information Act* in the case of an access to information request. For details with regard to personal information collected under this program, visit the Personal Information Bank; Health Canada; Health Products and Food Branch; Branch Incident Reporting System; HC PPU 088 at: http://infosource.gc.ca/inst/1476/1476-fedemp00-eng.asp

E. INVESTIGATION INFORMATION

1. **Investigative Actions and Timeline**

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This section only applies for preliminary & final, and final reports

2. **Root Cause of Problem**

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3. **Corrective Actions taken as a result of the investigation**

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