Republic of the Philippines
Department of Health
**FOOD AND DRUG ADMINISTRATION**

03 AUG 2018

**FDA ADVISORY**
No. 2018-240

**TO:** **THE GENERAL PUBLIC AND ALL CONCERNED HEALTHCARE PROFESSIONALS AND ESTABLISHMENTS**

**SUBJECT:** **Termination of the Voluntary Recall of AV-Set B DT INF-E Blood Tubing System with DVR No. 8574 (Article No. AP16641)**

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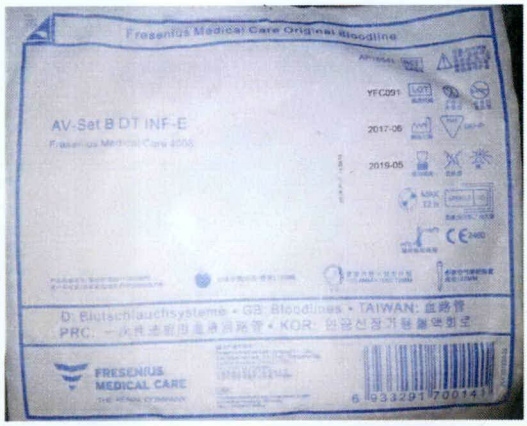
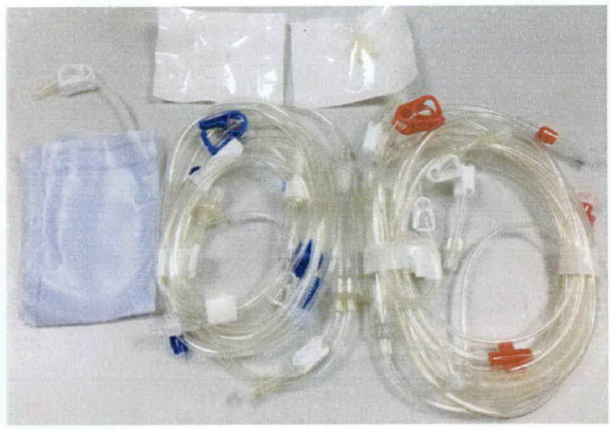
The Food and Drug Administration (FDA) informs the public that **Fresenius Medical Care Philippines, Inc.**, Marketing Authorization Holder (MAH), has reported that it has completed the recall and removal of **AV-Set B DT INF-E Blood Tubing System (see Figures 1 and 2) with DVR No. 8574 (Article No. AP16641)** from the Philippine market.



Figures 1 and 2: Photos of AV-Set B DT INF-E Blood Tubing System

Dissemination of the information to all concerned is requested.

**NELA CHARADE G. PUNO, RPh**
FDA Director General

20180424164745

Civic Drive, Filinvest City, Alabang 1781 Muntinlupa, Philippines
Trunk Line +63 2 857 1900 Fax +63 2 807 0751
Website: www.fda.gov.ph Email: info@fda.gov.ph