Republic of the Philippines
Department of Health
FOOD AND DRUG ADMINISTRATION

FDA ADVISORY
No. 2017-322

15 DEC 2017

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

SUBJECT: **TERMINATION OF THE VOLUNTARY PRODUCT RECALL OF COBRA DISTAL RADIUS FIXATOR**

The Food and Drug Administration (FDA) informs the public that **Orthopaedic International Inc.,** the Marketing Authorization Holder (MAH), has completed the recall, removal, and destruction of **Cobra Distal Radius Fixator** with **Lot Nos. 1601022 and 1600770** from the Philippine market.

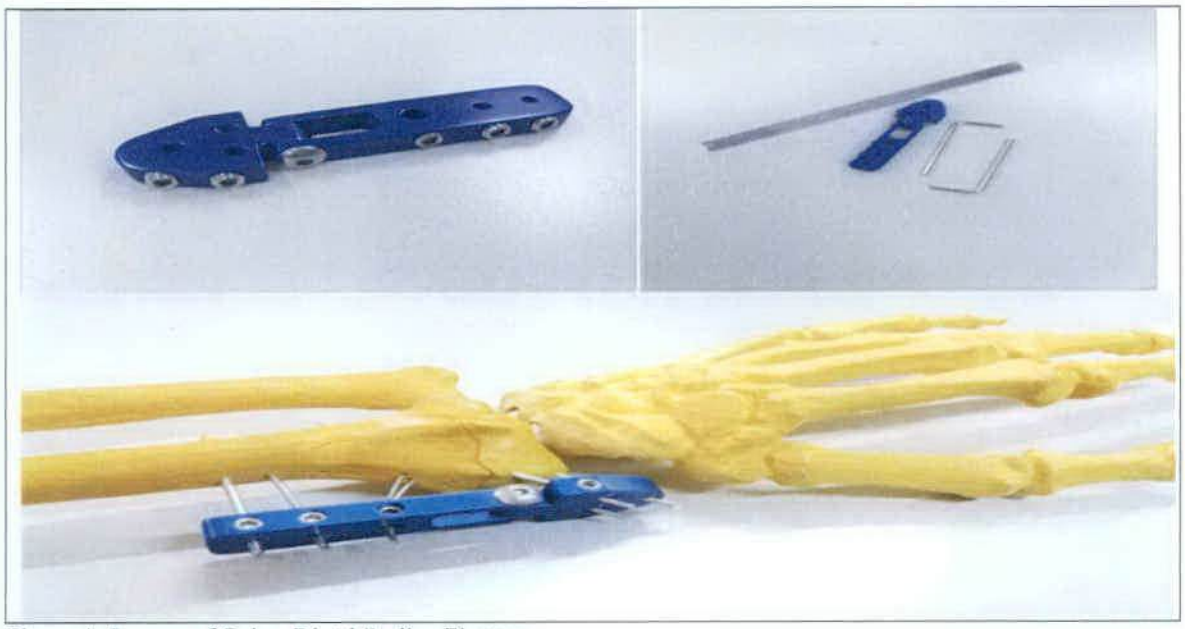

Figure 1: Images of Cobra Distal Radius Fixator

Dissemination of the above information to all concerned is requested.

NELA CHARADE G. PUNO, RPh
Director General

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