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Class 2 Device Recall Turon Impaction Fixture

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Class 2 Device Recall Turon Impaction Fixture

See Related Information

Date Initiated by Firm

February 09, 2017

Create Date

March 08, 2017

Recall Status<sup>1</sup>

Open<sup>3</sup>, Classified

Recall Number

Z-1412-2017

Recall Event ID

76456<sup>23</sup>

**Product Classification** 

Impactor<sup>24</sup> - Product Code HWA<sup>25</sup>

Product

Turon Impaction Fixture

**Code Information** 

101521L01, 115670L18, 128092L10, 136172L07, 136172L08, 141511L22, 37369L10A, 37369L10B, 37837L23A, 37837L23B, 37837L23C, 37837L23D, 51103L05E, 51103L05F, 52748L02A, 52748L02B, 52748L02C, 52748L02D, 52748L02E, 52748L02F, 52748L02G,

52748L02H, 57074L05A, 89910L10

Recalling Firm/ Manufacturer

Encore Medical, Lp 9800 Metric Blvd Austin TX 78758-5445

For Additional Information Contact Desiree Wells 512-832-9500

Manufacturer Reason

for Recall

During the Turon assembly, the impaction forces caused the polymer, black acetal copolymer from the Impaction Fixture to wear off on the lateral surface of the Humeral

Stem titanium plasma spray coating.

**FDA** Determined

Cause 2

Device Design

Action

There are two field safety notices, one for Consignees who have surgeons that use the impaction fixtures (Verson 1) and one for Consignees who had previously indicated their surgeons do NOT use the impaction fixtures (Verson 2). The two recall notification letters

were sent out on 2/9/17.

Quantity in Commerce

297 units

Distribution

US, South Korea, Australia, Canada, United Kingdom/Ireland, Germany

Total Product Life Cycle

TPLC Device Report<sup>26</sup>

updated as the status changes.

Links on this page:

<sup>&</sup>lt;sup>1</sup> A record in this database is created when a firm initiates a correction or removal action. The record is updated if the FDA identifies a violation and classifies the action as a recall, and it is updated for a final time when the recall is terminated. Learn more about medical device recalls<sup>27</sup>

<sup>&</sup>lt;sup>2</sup> Per FDA policy, recall cause determinations are subject to modification up to the point of termination of the recall. <sup>3</sup> The manufacturer has initiated the recall and not all products have been corrected or removed. This record will be