

Date: December 22, 2021

Subject: MEMO: AMT ENFit[®] + Legacy Product Offering

To Whom It May Concern,

AMT would like to remind our customers that we will continue to offer a complete Legacy Line and a complete ENFit[®] Line, pursuant to FDA regulations.

As of January 1, 2022 all GEDSA Manufacturer Members (AMT is not a member) will no longer manufacture transitions sets and adaptors sold separately from other devices, as stated in the July 1, 2021 *Revised ENFit*[®] *Connector Conversion Schedule*.

AMT is not a member of GEDSA and, while we believe in their efforts and align with their messaging of patient safety, our position has not changed.

AMT is a leading medical device manufacturing facility located in northeast Ohio. We make medical devices, but our business is people. We are committed to the highest level of quality, safety, and patient care. We are dedicated to being an active participant in our customers' lives as they make the transition to enteral nutrition. At AMT, we do not decide what devices or accessories our customers use, rather we allow them to choose.

GEDSA letter: <u>https://stayconnected.org/wp-</u> <u>content/uploads/2020/09/200914_REVISED_ENFit%C2%AE_Connector_Conversion_Schedule_U.</u> <u>S. Canada_Legacy_Connector_Production-Phase_Out_Dates-1-1.pdf</u>

Thank you for your continued business, and we welcome all feedback.

ENFit® is a registered trademark of GEDSA, Inc. or its affiliates.