

04 Dec 2020 | Analysis

Device Week, 4 December 2020 – MedWatch Question About Third-Party Servicers Slides Under Industry's Radar

by Shawn M. Schmitt

On this week's podcast: A change made by the US FDA to its MedWatch program that asks adverse event reporters whether a third party serviced a malfunctioning medical device went unnoticed by many in industry for nearly a year. We explain why, and tell how the servicer question will be helpful for manufacturers and the agency.

Listen to the podcast via the player below:

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Medtech Insight article addressing topics discussed in this episode:

• <u>Recent eMDR Change By FDA Asks Adverse Event Reporters About Third-Party Servicers</u>