MEDTECH INSIGHT
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06 Dec 2019 | **Analysis**

QUOTED. 6 December 2019. Jon Speer.

by

Medical device regulations that were set in place by the US FDA in the 1990s are evolving with the emergence of digital-health products and their iterative nature, medtech consultant Jon Speer says. Check out his comment here.

“We have to change the perception of what regulation is and isn’t.” – Jon Speer, founder, Greenlight Guru

- Find out more: *Tech Companies Grappling With A Brave New World Of Regulations While Developing Digital-Health Products*

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