MEDTECH INSIGHT
CITELINE COMMERCIAL

21 Jun 2018 | **News**

QUOTED. June 21, 2018. Bassil Akra.

by

There is confusion over whether it is appropriate for notified bodies to conduct “mock audits” in advance of the new EU Medical Device Regulations. Notified body executive Bassil Akra explains one reason why they can be very helpful here.

"The new regulations are not an upgrade – they are hugely disruptive, and preparing for them is like never having been on the market before.” –Bassil Akra, VP of global focus teams, TÜV SÜD Product Service

- Find out more: *Mock Audits: Has Ambiguity In EU Led To Unfair Advantage?*

Click here for a free trial of *Medtech Insight*