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QUOTED. Feb. 22, 2018. Dan Vukelich.

by

Check out what Association of Medical Device Reprocessors President Dan Vukelich had to say about his expectations in the European market now that new rules of the road are being established for reprocessing of single-use devices in the EU.

“We expect most countries will ultimately opt in to allow commercially available, CE-marked remanufactured single-use devices (SUDs) because such products are now held to the same safety and efficacy standards as all other medical devices.” –Dan Vukelich, president, Association of Medical Device Reprocessors

- Find out more: *Could New EU Regulatory Hurdles Put Up Barriers For Would-Be Reprocessors?*

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