Lean RAQA Systems, LLC to Advise FDA on Scalable Solutions for Good Manufacturing Practices

FDA confirms Michelle Lott from Lean RAQA Systems, LLC as the newest member of FDA Device Good Manufacturing Practices Advisory Committee.

TUCSON, AZ, USA, August 29, 2016 /EINPresswire.com/ -- Michelle Lott, Principal and Founder of Lean RAQA Systems, LLC, was recently confirmed as the Food and Drug Administration (FDA)'s newest member of the Device Good Manufacturing Practices (DGMP) advisory committee. The DGMP advisory committee reviews proposed regulations for good manufacturing practices governing the methods used in, and the facilities and controls used for, the manufacture, packing, storage, and installation of devices, and makes recommendations on the feasibility and reasonableness of the proposed regulations. Ms. Lott is one of two Industry Representatives on the committee of ten public, health professionals, and government representatives. Passionate about advocating for innovators and small medical device companies, Ms. Lott looks forward to advising FDA on scalable solutions for the medical device industry. Lean RAQA Systems, LLC specializes in regulatory and quality strategy.

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