



Need help with Establishment Registration and Medical Device Listing?

Global Regulatory Partners helps medical devices companies register their establishment and list their medical devices with FDA. To learn more about the process, check out our white paper on this subject or Contact Us today!



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Need help with US Registration?

Global Regulatory Partners helps medical devices companies register their products with the FDA and access U.S. market by providing the following services:

- Medical Device Classification
- Pre-Submission meetings with the FDA
- Medical Device Registration
- FDA 21 CFR Part 820 - Quality Management System Implementation



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