



Have you ever left a conference and realized that you forgot to ask the question you came to get answered? Ever wonder what the FDA's criteria may be when reviewing your PMA/510(k) submission? Join other at [MDMA's FDA Forum](#) on March 12-13, in Palo Alto, CA to get your questions answered.

[MDMA's 2018 FDA Forum](#) continues to grow as the conference to engage with leading policy makers and experts. Be sure not to miss out hearing from **EIGHT FDA officials** on how to get your technologies into the marketplace.

Confirmed FDA Speakers Include:

- **William Maisel**, Deputy Center Director for Science & Chief Scientist, CDRH, FDA
- **Sean Boyd**, Deputy Director for Regulatory Affairs, Office of Compliance, FDA
- **Abiy Desta**, Ombudsman, CDRH, FDA (via video conferencing)
- **Elizabeth Hillebrenner**, Associate Director for Programs and Performance, Office of In Vitro Diagnostics and Radiological Health, CDRH, FDA
- **Joshua Nipper**, Chief of Premarket Approval Staff, Office of Device

- Evaluation, CDRH, FDA
- **Bakul Patel**, Associate Director for Digital Health, Office of the Center Director, CDRH, FDA
 - **Marjorie Shulman**, Chief of Premarket Notification (510(k)) Section, Office of Device Evaluation, CDRH, FDA
 - **Suzanne Schwartz, MD, MBA**, Associate Director for Science and Strategic Partnerships, Office of the Center Director, CDRH, FDA

This is a rare opportunity to share your concerns and ask questions from some of the FDA's top officials.

MDMA Members - \$595

State Member - \$795

Non-Members - \$895

NCBIO Members get at \$100 discount.

To get that discount, contact Amber Niebauer, aniebauer@ncbioscience.net.

For more information and to register, [CLICK HERE](#)

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