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LCDR Neil A. Mafnas is a Senior Regulatory Operations Officer for the Medical Device Single Audit (MDSAP) Team in the Center for Devices and Radiological Health, Office of Compliance, Division of Import Compliance Operations. As part of the MDSAP Team, LCDR Mafnas' duties focus on IT project management, training and outreach for FDA staff, and as an MDSAP CDRH liaison for FDA's Office of Regulatory Affairs.

LCDR Mafnas started his FDA career in 2010, working as a Consumer Safety Officer for the General Hospital Devices Branch within CDRH, Office of Compliance, Division of Enforcement A. From there he transitioned to the Respiratory, ENT, General Hospital and Ophthalmic Devices Branch in the Division of Manufacturing and Quality of the Office of Compliance. In these positions, LCDR Mafnas conducted Quality System (QS) regulation reviews of Establishment Inspection Reports (EIRs), classified Medical Device Recalls, led Health Risk Assessments of device failures, investigated and processed medical device complaints, assisted with regulatory cases and provided inspectional guidance to FDA District Offices.

LCDR Mafnas spent just over six years as an active duty U.S. Air Force Logistics Officer. While in the Air Force, he held maintenance and engineering positions in: Andersen Air Force Base, Guam; Cape Canaveral Air Force Station, Florida; and Beale Air Force Base, California.

LCDR Mafnas received a Bachelor of Science degree in Biology from the University of Texas at Arlington and a Master of Science degree in Health Sciences from the University of Central Florida.