Committee on Energy and Commerce
U.S. House of Representatives

Witness Disclosure Requirement - "Truth in Testimony"
Required by House Rule XI, Clause 2(g)(5)

1. Your Name: Diane Wurzburger		
2. Your Title: Executive Regulatory Affairs, Americas & Global Strategic Policy GE Healthcare		
3. The Entity(ies) You are Representing: Medical Imaging and Technology Alliance		
4. Are you testifying on behalf of the Federal, or a State or local government entity?	**Yes**	**No** X
5. Please list any Federal grants or contracts, or contracts or payments originating with a foreign government, that you or the entity(ies) you represent have received on or after January 1, 2015. Only grants, contracts, or payments related to the subject matter of the hearing must be listed. None		
6. Please attach your curriculum vitae to your completed disclosure form.		

Signature: [redacted] **Date:** 3/27/17

Diane Wurzburger, JD RAC
Executive, Regulatory Affairs
GE Healthcare

Diane Wurzburger, J.D., RAC is currently Executive of Regulatory Affairs for GE Healthcare with responsibility for the United States, Canada and Latin America as well as Global Strategic Regulatory Policy. Diane has over 25 years of experience in the medical device industry, with responsibilities that have included strategic regulatory, quality system, clinical, compliance and reimbursement leadership for Class II and Class III medical devices. Diane is an active MITA member, serving on the Board of Directors as well as Chair of the Technical and Regulatory Committee. She was a MITA industry representative to the MDUFA IV negotiations with FDA. She is also a member of RAPS and is RAC certified. Diane holds a Bachelor of Science degree in Biomedical Engineering from Boston University and a Juris Doctor degree from Seton Hall Law School. She is a member of the New Jersey State Bar Association.