Industry Representative Pool for the Interests of the Medical Device Industry
Patient Engagement Advisory Committee

The FDA published a notice in the *Federal Register* (February 13, 2020, Vol. 85, No. 30, pp. 8298-8299) requesting that interested industry organizations select a pool of individuals, with varying areas of expertise, to represent industry interest for the Committee. The Commissioner or designee has the authority to select an individual from this pool to serve temporarily as a nonvoting member, depending on the meeting topic. In accordance with the process described in the notice, the following candidates were identified as participants in the pool, with terms continuing to April 30, 2025.

Industry Segment	Primary	Alternate
Cybersecurity	**Pat Baird** Senior Regulatory Specialist, Head of Global Software Standards, Philips	**Damien McPhillips, PMP, CSQE** Sr. Quality Manager, Global Software and Digital Health Boston Scientific Corp.
Communication of Benefit & Risk Information to Patients; Medical Device Labeling	**Jijo James, M.D., M.P.H.** Chief Medical Officer, Johnson and Johnson Medical Devices Companies & Global External Innovation	**Elisabeth George, MS** Elisabeth George Consulting LLC 7901 4th Street N. Suite 300 St. Petersburg, FL 33702
Health of Women/Pediatrics (Vulnerable Population Groups)	**Rose E. Blackburne, M.D., M.B.A.** Vice President, Global Head General Medicine Global Product Development	**Elisabeth George, MS** Elisabeth George Consulting LLC 7901 4th Street N. Suite 300 St. Petersburg, FL 33702
Patient Engagement	**Dan P. Stephens, Ph.D.** Principal Clinical Specialist, Global Advocacy Boston Scientific	**Elisabeth George, MS** Elisabeth George Consulting LLC 7901 4th Street N. Suite 300 St. Petersburg, FL 33702
Patient Preference Elicitation	**Dan P. Stephen, Ph.D.** Principal Clinical Specialist, Global Advocacy Boston Scientific	**Rose E. Blackburne, M.D., M.B.A.** Vice President, Global Head General Medicine Global Product Development
Patient-reported Outcomes Development, Validation and Use in Regulatory Studies or Clinical Practice	**David Amor.** Head of Quality and Regulatory Affairs Apple, Inc.	**Dan P. Stephens, Ph.D.** Principal Clinical Specialist, Global Advocacy Boston Scientific
Postmarket Studies, including observational and Registry-based Studies	**Elisabeth George, MS** Elisabeth George Consulting LLC 7901 4th Street N. Suite 300 St. Petersburg, FL 33702	**Jijo James, M.D.** Chief Medical Officer, Johnson and Johnson Medical Devices Companies & Global External Innovation
Digital Health Technology/Artificial Intelligence	**Diane M. Johnson** Senior Director North American Policy Global Digital Health Policy Lead, Johnson & Johnson	**Pat Baird** Senior Regulatory Specialist, Head of Global Software Standards, Philips