AGENDA

From 8.00am	**Networking Breakfast**
9:30-9.35am	**Opening Remarks** Robbie Walsh, Director, Regulatory Affairs, Boston Scientific
9:35-9:45am	**Overview of IMA Advocacy & Outreach** Emer Sherry, Senior Executive, Ibec
9:45-10:30am	**Quality Management Systems Impacts and Audit Task Force** Fidelma Sheerin, Chief Specialist, Corporate Quality, Stryker Ann Hunt, Snr Quality Systems Manager, Medtronic **Q&A**
10:30-11:00am	**Networking Tea & Coffee Break**
11:00-12:00pm	**HPRA Update** Nicola Hickie, Regulatory & Policy Manager, Medical Devices **Q&A** Nicola Hickie and Niall MacAleenan, Director of Medical Devices, HPRA
12:00-12:45pm	**Technical Documentation Task Force** Paul Cahalan, Regulatory Manager, Boston Scientific Ellen Looby, Senior Regulatory Affairs Specialist, Merit Medical Thomas Flannelly, Staff Engineer, Regulatory Affairs Stryker Elizabeth Delahunty, Associate Director, Regulatory Affairs, BD **Q&A**
12:45-2:00pm	**Networking Lunch**
2:00-2:30pm	**Regulatory Landscape for the UK (UKCA) and Switzerland (CH-REP)** Lynn Heaver, Regulatory Affairs Lead UK/Ireland, Johnson & Johnson
2:30-3:15pm	**Clinical Evaluation Task Force** Simon Rowan, Clinical Affairs Manager, Aerogen Joanna McCarthy, Associate Clinical Research Manager, Stryker Alice Forde, Director, Regulatory Affairs Europe, Medical Devices at Bausch + Lomb **Q&A**
3:15-3:20pm	**Closing Remarks** Global Access 2022, 12th & 13th October 2022, The Galmont Hotel, Galway
3:30pm	**Forum Close**