International Regulatory Developments

Regulatory requirements have been increasing globally in the last years as new jurisdictions develop their own regulatory systems. In such an environment, facilitating regulatory convergence and ensuring that the requirements being deployed around the world are in line with regulatory best practices is an essential part of the activities of MedTech Europe.

The trend where we see more and more jurisdictions and oversight stepping in to regulate medical technologies is likely to continue as the importance and value of medical technologies to health systems as a whole becomes widely recognized.