

# Regulation in far from Regular Times

## ABHI Webinar

27 May 2020, 14:00 - 16:30

| Time  | Topic   | Speaker  |
|-------|---|--|
| 14:00 | Introductions                                       | <ul style="list-style-type: none"><li>• <b>Phil Brown</b>, Director, Regulatory and Compliance, ABHI</li></ul>   |
| 14:15 | MDR Delay: What does it mean?                       | <ul style="list-style-type: none"><li>• <b>Dr Camilla Fleetcroft</b> - Group Manager – Devices Regulatory Affairs, MHRA</li><li>• <b>Oliver Bisazza</b> - Director Regulations &amp; Industrial Policy, MedTech Europe</li><li>• <b>Shuna Mason</b>, Partner, CMS Cameron McKenna Nabarro Olswang LLP</li></ul> <p>Q&amp;A chaired by Phil Brown</p>                       |
| 15:00 | Fit for a pandemic: Humanitarian use of regulations | <ul style="list-style-type: none"><li>• <b>Michael Kipping</b>, Innovation Lead - Biomedical Catalyst, Innovate UK &amp; currently seconded to MHRA</li><li>• <b>Steve Lee</b>, Senior Regulatory Policy Advisor - IVDR, MHRA</li><li>• <b>James Pink</b>, Executive Vice President, Consulting Services, NSF International</li></ul> <p>Q&amp;A chaired by Phil Brown</p> |
| 15:45 | What next? Regulation in a Global world             | <ul style="list-style-type: none"><li>• <b>Joe Gatewood</b>, Vice President, Global Strategy and Analysis, AdvaMed</li><li>• <b>Lynn Heaver</b>, Regulatory Affairs Lead UK&amp;I, Johnson &amp; Johnson</li></ul> <p>Q&amp;A chaired by Phil Brown</p>  |
| 16:15 | Wrap up   | <ul style="list-style-type: none"><li>• <b>Phil Brown</b>, Director, Regulatory and Compliance, ABHI</li></ul>   |