

## MDSAP Regulatory Authority Contact Information

<p style="text-align: center;"><b>AUSTRALIA – TGA</b></p> <p style="text-align: center;"><a href="mailto:MDSAP@tga.gov.au">MDSAP@tga.gov.au</a></p>	<ul style="list-style-type: none"> <li>• Medical Device Single Audit Program (MDSAP) Medical Devices Branch (MDB) Therapeutic Goods Administration (TGA) Department of Health PO Box 100, Woden, ACT 2606 Australia</li> </ul>
<p style="text-align: center;"><b>BRAZIL – ANVISA</b></p> <p style="text-align: center;"><a href="mailto:MDSAP.ATENDIMENTO@ANVISA.GOV.BR">MDSAP.ATENDIMENTO@ANVISA.GOV.BR</a></p>	<ul style="list-style-type: none"> <li>• Anvisa – Agência Nacional de Vigilância Sanitária Setor de Indústria e Abastecimento (SIA) - Trecho 5, Área Especial 57 / Lote 200 Brasília (DF) – Brazil POSTAL CODE: 71205-050</li> </ul>
<p style="text-align: center;"><b>CANADA – HC</b></p> <p style="text-align: center;"><a href="mailto:QS_MDB_HC@hc-sc.gc.ca">QS_MDB_HC@hc-sc.gc.ca</a></p>	<ul style="list-style-type: none"> <li>• Quality Systems Division Medical Devices Bureau 2934 Baseline Road Qualicum, Tower B Ottawa, Ontario, Canada K1A 0K9 Address Locator: 3403A</li> </ul>
<p style="text-align: center;"><b>JAPAN – PMDA</b></p> <p style="text-align: center;"><a href="mailto:MDSAP@pmda.go.jp">MDSAP@pmda.go.jp</a></p>	<ul style="list-style-type: none"> <li>• Japan’s Ministry of Health, Labour and Welfare Ministry of Health, Labour and Welfare Medical Device Evaluation Division, Pharmaceutical Safety and Environmental Health Bureau 1-2-2, Kasumigaseki, Chiyoda-ku, Tokyo 1008616 Japan</li> <li>• Pharmaceuticals and Medical Devices Agency Office of Manufacturing/Quality and Compliance- Division of Registered Certification Body Assessment Shin-kasumigaseki Bldg. 3-3-2 Kasumigaseki, Chiyoda-ku, Tokyo 100-0013 Japan</li> </ul>
<p style="text-align: center;"><b>US – FDA</b></p> <p style="text-align: center;"><a href="mailto:MDSAP@FDA.HHS.GOV">MDSAP@FDA.HHS.GOV</a></p>	<ul style="list-style-type: none"> <li>• US FDA – Food and Drug Administration Center for Devices and Radiological Health Office of Compliance – Division of International Compliance Operations – Medical Device Single Audit Pilot Program (MDSAP) 10903 New Hampshire Avenue Building 66 Room 3543 Silver Spring, MD 20993-0002 Tel.: 301-796-5500 (main number)</li> </ul>