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ASAP INTERNATIONAL SDN. BHD. (1157980-X)

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EU DECLARATION of CONFORMITY	
Manufacturer:	<b>ASAP International Sdn. Bhd.</b> No. 1, Jalan Sitar 33/6, Seksyen 33, 40400 Shah Alam, Selangor, Malaysia. SRN: MY-MF-000004500
Medical Devices & Basic UDI- DI:	<ul> <li>Powdered Latex Examination Gloves (955589090ASAPPPL5E)</li> <li>Powder Free Latex Examination Gloves (955589090ASAPPFL4G)</li> <li>Powder Free Nitrile Examination Gloves (955589090ASAPPFN4L)</li> </ul>
Classification:	Class I Medical Devices according to Medical Device Regulation (EU) 2017/745, Annex VIII
Harmonised Standards:	EN455-1:2000, EN 455-2:2015, EN455-3:2015, EN455-4:2009
<b>EC REP</b> European Authorized Representative:	ASAP INNOVATIONS LIMITED 7 Saggart Lakes, Saggart, Dublin, D24 PY01, Ireland. SRN: IE-AR-000002548
This EU declaration of conformity is issued under the sole responsibility of manufacturer, ASAP International Sdn. Bhd. We hereby declare that device covered by the present declaration is in conformity with the Medical Device Regulation (EU) 2017/745 and with the above-mentioned standards. All supporting documentation is retained at the premises of the manufacturer.	
Place, Date of Declaration:	Malaysia., May 26, 2021
Signature:	John
	Mr. Chin Tze Weng Senior Global Business Development Manager

