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<b>CONFORMITY TO TYPE BASED ON QUALITY ASSURANCE OF THE PRODUCTION PROCESS</b>		<b>Certificate N° 16/US/3838</b> Index.3 – Rev.0	
<b>In accordance with the requirements of Module D of the « Pressure Equipment Directive » 2014/68/EU</b>			
<b>Manufacturer</b>	Name	<b>AIRSEP Corporation</b>	
	Address	260 Creekside Drive Buffalo, NY 14228 USA	
<b>Concerned Scope:</b> <p>Manufacturing, inspection and testing of oxygen generators including their components</p>			
<b>Certified products</b> as per module B certificate n°: <b>16/US/3920</b>			
<p>This is to certify that the Quality Management System of the above-mentioned manufacturer has been assessed against the requirements of annex III Module D of the Pressure Equipment Directive 2014/68/EU, conforming the requirements for equipment &amp; assemblies shown above. The approval is subject to the continued maintenance of the Quality System in accordance with the requirements of the above Directive, this shall be controlled by intermediate audits, inspections and surveys.</p>			
<p>The manufacturer is allowed to affix the “CE” mark followed with our notified body identification number 0029 to the approved equipment in the conditions described in the Directives.</p> <p style="text-align: center;"><b>The approval is valid until 3<sup>rd</sup> January 2025</b></p>			
Approval Date : 10 <sup>th</sup> January 2022	Name : B. Nève	Position	<b>General Manager</b>
Notified body identification number :	<b>0029</b>	Signature	
Notified body :	<b>APRAGAZ VZW/ASBL</b>		
Notified body reference :	<b>P16526/000</b>		