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NOTIFIED BODY EC CERTIFICATE OF CONFORMITY		Certificate N°																		
In accordance with Appendix II of the Medical Devices Directive 93/42/EC (excluding section 4)		05/FR/918-3-REV 0																		
Manufacturer	Name	AIR LIQUIDE S.A. – DMC - CRYOPAL																		
	Address	Parc Gustave Eiffel – Avenue Gutenberg, 8 Bussy-Saint-Georges FR – 77607 Marne-La-Vallée FRANCE																		
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Concerned Equipment : Category : Ila																				
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<p>This is to certify that the Quality Management System of the above mentioned manufacturer has been assessed against the requirements of Annex II of the Medical Devices Directive 93/42/EC and conforms to the requirements for the equipment 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 approved equipment in the conditions described in the Directive.</p> <p style="text-align: center;">The approval is valid until 16/06/2010.</p>																				
Date : 11/08/2008	Name : Ch. Leplat	Position General Manager																		
Notified body identification number :	0029	Signature 																		
Notified body stamp :	<p>APRAGAZ Belgium Inspecting Authority</p>																			
Notified body reference :	0502/P7338/ZMEDI01																			