By Royal Charter

EC Design-Examination Certificate

Directive 93/42/EEC on Medical Devices, Annex II Section 4

No. **CE 572880**

Issued To: **Preservation Solutions, Inc.**
1099 Proctor Drive
Elkhorn
Wisconsin
53121
USA

In respect of:

CoStorSol and MaPerSol sterile solution containing Adenosine, Allopurinol and Mannitol

BSI has performed a design examination on the above devices in accordance with the Council Directive 93/42/EEC, Annex II Section 4. The design conforms to the requirements of this directive. For marketing of these products an additional Annex II excluding Section 4 certificate is required.

For and on behalf of BSI, a Notified Body for the above Directive (Notified Body Number 0086):

Frank Lee, EMEA Compliance & Risk Director

First Issued: **30 July 2012** Date: **31 May 2016** Expiry Date: **29 July 2017**

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Supplementary Information to CE 572880

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Product	Container Volume	Catalog/Part Number
CoStorSol®	500 mL	PS006CE
CoStorSol®	1000 mL	PS004CE
CoStorSol®	2000 mL	PS007CE
MaPerSol®	1000 mL	PS005CE

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Certificate History

Date	Reference Number	Action
30 July 2012	10123172	First issue.
27 March 2015	10151632 10141213	Change of address. Addition of 2000mL and 500mL CoStorSol variants. Updated part numbers.
31 May 2016	10159839	Shelf life extension to 18 months for CoStorSol 500mL, 1L and 2L and MaPerSol 1L.

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