

CERTIFICATE OF REGISTRATION

This is to certify that the management system of:

Mediplast AB

Main site: Bronsåldersgatan 2, SE-213 76 Malmö, Sweden
Box 1004, SE-212 10 Malmö, Sweden

Additional sites according to appendix

has been registered by Intertek as conforming to the requirements of

SS-EN ISO 13485:2016

The management system is applicable to:

Design, development, and manufacturing of disposable, non-active sterile and non-sterile medical devices and distribution of medical devices primarily to Healthcare sector.

Repair, calibration and preventive maintenance of non-active, non-implantable and general active medical devices

*Previously certified by Intertek Certification AB to date 9 November 2020

Certificate Number:

0107192

Revision:

03

Initial Certification Date:

10 June 1998*

Date of Certification Decision:

11 October 2024

Certificate Valid Date:

11 October 2024

Certificate Expiry Date:

26 June 2027



Ackred. nr. 10426
Certifiering av ledningssystem
ISO/IEC 17021-1



Intertek

Mikael Hagelin

Certification Authority, Intertek Medical
Notified Body AB

Intertek Medical Notified Body AB
P.O. Box 1103, SE-164 22 Kista, Sweden



APPENDIX

This appendix identifies the locations by the management system of

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Mediplast AB

This appendix is linked to certificate #0107192-03 and cannot be shown nor reproduced without it.

Hospidana ApS

Kleins Vej 6, 4930 Maribo, Denmark

manufacturing of disposable non-active non-sterile medical devices, customer communication sales and distribution of medical devices.

Mediplast A/S

Marielundvej 46E, DK-2730 Herlev, Denmark

Customer communication and sales of medical devices.

Mediplast AS

Bjørnstjerne Bjørnsonsgate 110, 3044 Drammen, Norway

Customer communication and sales of medical devices.

Mediplast Fenno Oy

Äyritie 12 B, Vantaa, 01510, Finland

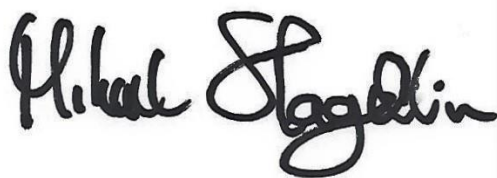
Customer communication and sales of medical devices.



APPENDIX

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