

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

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

Certificate Number:

0107192-01

Initial Certification Date:

10 June 1998*

Date of Certification Decision:

25 May 2021

Certificate Valid Date:

27 June 2021

Certificate Expiry Date:

26 June 2024



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



Intertek

Brian Mather

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

Page 1

Mediplast AB

This appendix is linked to certificate #0107192-01 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

Tollbugatan 115, NO-3041 Drammen, Norway

Customer communication and sales of medical devices.



APPENDIX

Certificate Number: 0107192-01
Initial Certification Date: 10 July 1998
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Page 2



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