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CERTIFICATE OF REGISTRATION

Certificate Number:
0107192

Revision:
02

Initial Certification Date:
10 June 1998*

Date of Certification Decision:
17 June 2024

Certificate Valid Date:
27 June 2024

Certificate Expiry Date:
26 June 2027

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

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

ISO 13485:2016 CERTIFICATION
Intertek

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.

Brian Mather
Certification Authority, Intertek Medical Notified Body AB

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

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

APPENDIX

This appendix identifies the locations by the management system of

Mediplast AB

This appendix is linked to certificate #0107192-02 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.
Mediplast Fenno Oy Äyritie 12 B, Vantaa, 01510, Finland	Customer communication and sales of medical devices.

APPENDIX

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