

# **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

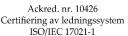
**Certificate Valid Date:** 

27 June 2021

**Certificate Expiry Date:** 

26 June 2024









#### **Brian Mather**

Certification Authority, Intertek Medical Notified Body AB

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







## **APPENDIX**

This appendix identifies the locations by the management system of

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### **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**

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