

ConvaTec Limited
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Declaration for Use of the Extended Transitional Period Article 120 - Regulation (EU) 2017/745

Article 120 in Regulation (EU) 2017/745 has been amended by Regulation (EU) 2023/607 to extend the transitional period for legacy devices beyond 26 May 2024, providing certain conditions are fulfilled.

This letter confirms that the products listed in Appendix 1 were covered by a valid certificate issued by a Notified Body (BSI The Netherlands; 2797) in accordance with Directive 93/42/EEC after 20 March 2023. The listed products are therefore eligible to avail of the extended transitional periods as per Article 120(3a) in Regulation (EU) 2017/745, as amended.

For the extended transitional period to apply, the following conditions stated in Article 120(3c) shall be met:

- (a) those devices continue to comply with Directive 90/385/EEC or Directive 93/42/EEC, as applicable;
- (b) there are no significant changes in the design and intended purpose;
- (c) the devices do not present an unacceptable risk to the health or safety of patients, users or other persons, or to other aspects of the protection of public health;
- (d) no later than 26 May 2024, the manufacturer has put in place a quality management system in accordance with Article 10(9);
- (e) no later than 26 May 2024, the manufacturer or the authorised representative has lodged a formal application with a notified body in accordance with Section 4.3, first subparagraph, of Annex VII for conformity assessment in respect of a device referred to in paragraph 3a or 3b of this Article or in respect of a device intended to substitute that device, and, no later than 26 September 2024, the notified body and the manufacturer have signed a written agreement in accordance with Section 4.3, second subparagraph, of Annex VII.

As Legal Manufacturer, we - ConvaTec Limited, First Avenue, Deeside Industrial Park, Deeside, Flintshire, CH5 2NU, United Kingdom (SRN; GB-MF-000001770), - herby declare, under our sole responsibility, that the conditions for the extension of the transitional period as stated in Article 120(3c) in Regulation (EU) 2017/745, as amended by Regulation (EU) 2023/607, are fulfilled. The products listed in Appendix 1 can thereby be placed on the market or put into service until the following dates:

| Class III and class IIb implantable devices: | 31 December 2027. |
|---|-------------------|
| Class IIb (non-implantable), class IIa and class I (sterile/measuring) devices: | 31 December 2028. |

DocuSigned by:

Signer Name: Steeve Lamvohee
Signing Reason: I approve this document
Signing Time: Jun 30, 2023 | 8:07:32 PM BST
79126EF62CD04B7AA0479B4D5AA7AE98

Jun 30, 2023

Steeve Lamovohee

Issue Date

Head of IC/AWC Regulatory Affairs



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Appendix 1: Product List

| Device Name | Basic UDI | MDR Device Classification | Transitional Period End Date | MDD Certificate Refernece | MDD Certificate Expiry Date | Notified Body# |
|---------------------------------|-----------------|-----------------------------|---------------------------------|------------------------------|--------------------------------|-------------------|
| Aquacel Surgical Cover Dressing | 768455AWC00203C | Class III | 31/12/2027 | CE 00364 CE 00380 | 20/10/2023 26/05/2024 | 2797 |
| Duoderm Extra Thin | 768455AWC00303F | Class III | 31/12/2027 | CE 00364 CE 00380 | 20/10/2023 26/05/2024 | 2797 |
| Duoderm Signal | 768455AWC00313H | Class III | 31/12/2027 | CE 00364 CE 00380 | 20/10/2023 26/05/2024 | 2797 2797 |
| Duoderm CGF | 768455AWC00323K | Class III | 31/12/2027 | CE 00364 CE 00380 | 20/10/2023 26/05/2024 | 2797 2797 |
| Duoderm CGF Border | 768455AWC00293W | Class III | 31/12/2027 | CE 00364 CE 00380 | 20/10/2023 26/05/2024 | 2797 2797 |
| Aquacel Foam | 768455AWC00063J | Class Ilb (non-implantable) | 31/12/2028 | CE 00364 | 20/10/2023 | 2797 |
| Aquacel Foam Pro | 768455AWC00073L | Class IIb (non-implantable) | 31/12/2028 | CE 00364 | 20/10/2023 | 2797 |
| Aquacel Extra & WSF | 768455AWC00213E | Class IIb (non-implantable) | 31/12/2028 | CE 00364 | 20/10/2023 | 2797 |
| Carboflex | 768455AWC00083N | Class IIb (non-implantable) | 31/12/2028 | CE 00364 | 20/10/2023 | 2797 |
| Duoderm Gel | 768455AWC00093Q | Class IIb (non-implantable) | 31/12/2028 | CE 00364 | 20/10/2023 | 2797 |
| ConvaTec Foam Lite | 768455AWC00193T | Class IIb (non-implantable) | 31/12/2028 | CE 00364 | 20/10/2023 | 2797 |
| Kaltostat | 768455AWC00053G | Class IIb (non-implantable) | 31/12/2028 | CE 00364 | 20/10/2023 | 2797 |
| Avelle NPWT Dressing | 768455AWC00033C | Class IIb (non-implantable) | 31/12/2028 | CE 00364 | 20/10/2023 | 2797 |
| Avelle NPWT Pump | 768455AWC00023A | Class IIa | 31/12/2028 | CE 00364 | 20/10/2023 | 2797 |
| Hydrocolloid Sealing Strips | 768455AWC00163M | Class Is | 31/12/2028 | CE 56172 | 20/10/2023 | 2797 |
| Loop Ostomy Rod | 7684550ST0005F5 | Class IIa | 31/12/2028 | CE 00364 | 20/10/2023 | 2797 |

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