

### EC-Conformity Declaration

(according to Appendix IX of the Medical Device Regulation (EU) 2017/745 of the European Parliament and of the Council)

We,  
Bauerfeind AG  
Triebeser Str. 16  
D-07937 Zeulenroda-Triebes  
SRN: ./.

declare in exclusive accountability, that our products:

Name	Intended Use	Basis-UDI-DI
VenoTrain® business	Compression stockings for the long-term therapy of phlebological and / or lymphological disorders as well as, in special cases, for the long-term therapy of lipedema	4046445BUS00000000XA
VenoTrain® look	Compression stockings for the long-term therapy of phlebological and / or lymphological disorders as well as, in special cases, for the long-term therapy of lipedema	4046445LOO000000000YN
VenoTrain® clinic	Compression stockings for the long-term therapy of phlebological and / or lymphological disorders as well as, in special cases, for the long-term therapy of lipedema	4046445CLI000000000GW
VenoTrain® cocoon	Compression stockings for the long-term therapy of phlebological and / or lymphological disorders	4046445COC000000000FB
VenoTrain® delight/ VenoTrain® delight T	Compression stockings for the long-term therapy of phlebological and / or lymphological disorders as well as, in special cases, for the long-term therapy of lipedema	4046445DEL000000000DP
VenoTrain® discrétion	Compression stockings for the long-term therapy of phlebological and / or lymphological disorders	4046445DIS000000000NC
VenoTrain® impuls	Compression stockings for the long-term therapy of phlebological and / or lymphological disorders as well as, in special cases, for the long-term therapy of lipedema	4046445IMP000000000UN

#### ADDRESS

Bauerfeind AG  
Triebeser Straße 16  
07937 Zeulenroda-Triebes  
Germany

#### CONTACT

P +49(0) 36628 66-1000  
F +49(0) 36628 66-1999  
E info@bauerfeind.com

#### DELIVERY ADDRESS

Weißendorfer Straße 5  
07937 Zeulenroda-Triebes  
Germany

Bauerfeind AG  
AG Jena HRB 206561

#### CHIEF EXECUTIVE OFFICER

Rainer Berthan

#### MEMBER OF THE EXECUTIVE BOARD

Andreas Lauth

#### CHAIRMAN OF THE BOARD

Prof. Hans B. Bauerfeind

[BAUERFEIND.COM](http://BAUERFEIND.COM)

VenoTrain® micro	Compression stockings for the long-term therapy of phlebological and / or lymphological disorders as well as, in special cases, for the long-term therapy of lipedema	4046445MIC00000000KK
VenoTrain® pure	Compression stockings for the long-term therapy of phlebological and / or lymphological disorders as well as, in special cases, for the long-term therapy of lipedema	4046445VTP00000000FP
VenoTrain® soft	Compression stockings for the long-term therapy of phlebological and / or lymphological disorders as well as, in special cases, for the long-term therapy of lipedema	4046445SOF00000000YY
VenoTrain® soft S	Compression stockings for the long-term therapy of phlebological and / or lymphological disorders	4046445SOFs000000005D
VenoTrain® ulcertec	Compression stockings for the long-term treatment of severe phlebological disorders in the legs	4046445ULC00000000W2
VenoTrain® curafLOW	Compression stockings for the long-term therapy of lymphological disorders as well as for the long-term therapy of lipedema	4046445CUR00000000XJ
VenoTrain® angioflow	Long-term treatment of chronic venous insufficiency (CVI) in the case of initial PAOD, i.e. with an ankle-brachial index <sup>1</sup> of between 0.5 and 0.9 and an arterial pressure of 60 mmHg at the ankle	4046445ANG00000000FC
VenoTrain® glider	accessory to a medical device. It makes it easier to put on medical compression stockings for the lower and upper extremities	4046445GLI00000000LS
VenoTrain® glider plus	accessory to a medical aid. This product makes it easier to put on medical compression stockings for the lower extremities (up to compression class 3)	4046445GLIP000000007A

comply with the General Safety and Performance Requirements according to Annex I of the Medical Device Regulation (EU) 2017/745. Within the conformity evaluation process there has been no participation of a Notified Body.

The products have been classified according to Annex VIII, Chapter III, MDR 2017/745 in risk class I. They correspond to the given provisions of the Regulation (EU) 2017/745 for Medical Devices.

Zeulenroda-Triebes, 25.05.2021



Ines Exner  
Person Responsible for Regulatory  
Compliance  
Bauerfeind AG



Petra Schroeder  
Head of Regulatory Affairs Int.  
Bauerfeind AG