

Dichiarazione di Conformità

Noi come azienda produttrice

ERKA.Kallmeyer Medizintechnik GmbH & Co. KG

Im Farchet 15
83646 Bad Toelz
Germania

ERKA.
THE ORIGINAL
Made in Germany since 1889

dichiariamo in assoluta responsabilità, che i prodotti medicali:

Stetoscopi

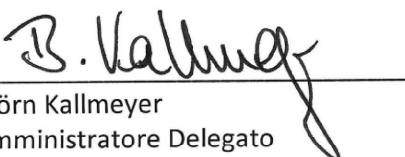
Artikel	Bezeichnung	UMDNS-Code	Klasse
520.	Finesse Light	13-755	I
520.	Finesse Light II		
521.	Finesse Light Child		
525.	Sensitive		
525.	Sensitive Porsche Design		
526.	Finesse ² light		
528.	Finesse light Simplex		
531.	Precise		
535.	Finesse ²		
536.	Finesse ² Child		
541.	Erkaphon Duo		
541.	Lehr-Erkaphon Duo		
542.	Erkaphon Duo Alu		
542.	Erkaphon Duo Black Line		
543.	Erkaphon flach		
543.	Lehr-Erkaphon flach		
544.	Erkaphon Alu-flach		
544.	Erkaphon Black Line		
545.	Erkaphon Duo Child		
547.	Erkaphon Child flach		
549.	Finesse Child		
550.	Finesse		
550.	Erkaphon Deluxe		
570.	Classic		

si attengono a tutte le disposizioni della Direttiva 93/42/CEE, Allegato VII

Organismo designato: DEKRA Certifications GmbH
Indirizzo: Handwerkstraße 15
D-70565 Stuttgart
Numero d'identificazione: 0124

La Dichiarazione di Conformità è valida fino al 29.11.2020

Bad Toelz, li 20.11.2017


Björn Kallmeyer
Amministratore Delegato

Declaration of conformity

We, as manufacturer

ERKA.Kallmeyer Medizintechnik GmbH & Co. KG

Im Farchet 15
83646 Bad Toelz
Germany

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hereby declare in sole responsibility that the medical devices:

Stethoscopes

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meet all the applicable provisions set out in the directive 93/42/EEC, annex VII.

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Bad Toelz, 20.11.2017


Björn Kallmeyer
Managing Director