

## EC-Declaration of Conformity on Medical Devices

According to annex II of Medical Device Directive 93/42/EEC (2007/47/EG)

Manufacturer:

**RECK-Technik GmbH & Co. KG** Engineering Division Sector  
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Name of Product:

Basic UDI-DI: 42 601 937 10018	<b>REF</b>	<b>SN</b>
<b>MOTomed viva2</b>	200.003	xxx-VM2-003xxx
<b>MOTomed viva2 stativ</b>	200.012	xxx-VM2-012xxx
<b>MOTomed viva2 light</b>	200.004	xxx-VM2-004xxx
<b>MOTomed viva2 light stativ</b>	200.021	xxx-VM2-021xxx
<b>MOTomed viva2 Parkinson</b>	200.008	xxx-VM2-008xxx
Basic UDI-DI: 42 601 937 10049	<b>REF</b>	<b>SN</b>
<b>MOTomed letto2</b>	279.003	xxx-LET2-003xxx
<b>MOTomed letto2</b>	279.024	xxx-LET2-024xxx
<b>MOTomed letto2</b>	279.008	xxx-LET2-008xxx
<b>MOTomed letto2</b>	279.016	xxx-LET2-016xxx
Basic UDI-DI: 42 601 937 10063	<b>REF</b>	<b>SN</b>
<b>MOTomed gracile12</b>	594.003	xxx-KIG12-003xxx
<b>MOTomed gracile12 ATAP</b>	594.003	xxx-KIG12-003xxx
	+	
	599.000	

Product options: all (according to current pricelist)

Product classification: IIa (rule 9, Medical Device Directive 93/42/EEC)

We hereby declare under our sole responsibility that the above products comply with the relevant provisions of the Medical Devices Directive 93/42/EEC (2007), Annex II, Section 3 and that we are solely responsible for issuing this declaration of conformity.

This Declaration of Conformity is valid until 02.08.2023 or until a revised declaration is made due to product modifications.

Following party was involved in the evaluation of the conformity:  
DEKRA Certification GmbH, Handwerkstrasse 15, 70565 Stuttgart, GERMANY  
Notified Body No. 0124



Betzenweiler, 03.08.2018

**Andreas Reck** Executive Director