



# EC-CERTIFICATE

(Full quality assurance system)



This is to certify that the company

## Ormed GmbH

Bötzingen Straße 90  
79111 Freiburg  
Germany

has implemented and maintains a full quality assurance system which applies to the products at every stage from design to final controls.

Through an audit, documented in a report, performed by DQS Medizinprodukte GmbH, it was verified that the management system fulfills the requirements of

## Annex II – excluding Section 4 of Council Directive 93/42/EEC concerning medical devices

with respect to the following medical devices:

Devices for passive joint mobilization (CPM-Therapy / Continuous Passive Motion) and Devices for active joint mobilization (CAM-Therapy / Controlled Active Motion) according annex.

The manufacturer is subject to surveillance according to Annex II, Section 5. The CE marking with the Notified Body Identification Number (0297) may be affixed on the devices listed in the certificate. An EC Design Examination Certificate according to Annex II, Section 4 is required for class III devices covered by this certificate. The certificate is in the case of class I(s) devices (I(s) = class I products placed on the market in sterile conditions) limited to the aspects of manufacture concerned with securing and maintaining sterile conditions. The certificate is in the case of class I(m) devices (I(m) = class I devices with a measuring function) limited to the aspects of manufacture concerned with the conformity of the products with the metrological requirements.

Certificate registration no.	020812 MR2
Certificate unique ID	170771100
Effective date	2020-08-07
Expiry date	2024-05-26
Frankfurt am Main	2020-08-07

## DQS Medizinprodukte GmbH

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DQS Medizinprodukte GmbH is a Notified Body according to Council Directive 93/42/EEC concerning medical devices with the Identification Number 0297.



**Annex to certificate**  
**Certificate registration No.: 020812 MR2**  
**Certificate unique ID: 170771100**  
**Effective date: 2020-08-07**



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<b>Device family</b>	<b>Products</b>	<b>UMDNS-Code</b>	<b>Class</b>
Rehabilitation devices	ARTROMOT®-K1	17-138	Ila
	ARTROMOT®-SP3	17-138	Ila
	ARTROMOT®-S3	17-139	Ila
	ARTROMOT®-E2	17-139	Ila
	ARTROMOT® ACTIVE-K	15-220	Ila
	ARTROMOT®-S4	17-139	Ila