

## EU DECLARATION OF CONFORMITY

**Name and address of the manufacturer:** Guangdong Yuehua Medical Instrument Factory Co., Ltd.  
/  
Rongsheng Science and Technology Zone, Daxue Road, 515063  
Shantou, PEOPLE'S REPUBLIC OF CHINA  
SRN: CN-MF-000004539

**EU Authorized Representative:/** Eunitor GmbH  
ADD: Kennedydamm, 5, Düsseldorf, 40476, Germany.  
SRN: DE-AR-000005081

As the manufacturer of the following medical device, we herewith declare under our sole responsibility that the stated medical device:

**Name of the medical device: /** Alternating Pressure Mattress

**Basic UDI-DI: /** 694474954001M4, 694474954005MC,  
694474950030LF, 694474954004MA, 694474954006ME,  
694474950047LY, 694474950082M2, 694474950053LT,  
694474950055LX

**Product model:/** QDC-303, P4000IIE(B), QDC-303+P4000IIE(B),  
QDC-300B, P4000IIE(C), QDC-300B+P4000IIE(C),  
QDC-5010E+P3000N2EB, QDC-8010+P3000A2QB3,  
QDC-8080+P3000A2QB3

**Product code:/** EMDN CODE: V080701, ACTIVE ANTI-DECUBITUS MEDICAL MATTRESSES

**Intended purpose: /** Prevent pressure sore in bedridden patients.

**Trade name:/**



**of class: /** Rule 1, Class I

is in conformity with Regulation (EU) 2017/745 and with any other relevant Union legislation that provides for the issuing of this EU Declaration of Conformity. The declaration is valid in connection with the "final inspection report" of the device. /

**Conformity assessment procedure:/** Declare the conformity of the abovementioned products by issuing this EU Declaration of Conformity after drawing up the technical documentation set out in Annexes II and III of Regulation (EU) 2017/745 /



**CS reference: /** None

2022.11.16  
Ort, Datum / Place, date /  
Lieu, date / Luogo, data

Leonlin  
Name und Funktion / Name and function /  
Nom et fonction / Nome e funzion