Number: 6118114CE01

EU Quality Management System Certificate

Conformity Assessment Regulation 2017/745 on Medical devices, Annex IX Chapter I and III

Manufacturer:

Yian Medical Technology (Haining) Co., Ltd.

1st Floor Area 1,2nd Floor Area 1, Building A, No. 2 Caohejing Road, Haining Economic Development Zone, Haichang Street, Haining City 314400 Jiaxing City, Zhejiang Province China

SRN ID.: CN-MF-000027269

DEKRA grants the right to use the EC Notified Body Identification Number illustrated below to accompany the CE Marking of Conformity on the products concerned conforming to the required Technical Documentation and meeting the provisions of the EU- Regulation which apply to them:

0344

Supplement to certificate: 6096257CN

Authorized Representative:

Shanghai International Holding Corp. GmbH (Europe)

Eiffestrasse 80, 20537 Hamburg, Germany

DEKRA hereby declares that the above mentioned manufacturer fulfils the relevant requirements of EU Regulation 2017/745, including all subsequent amendments for the above mentioned conformity assessment. The manufacturer authorized representative is subject to periodic surveillance as required for the applicable conformity assessment in accordance to Regulation 2017/745.

DEKRA Certification B.V.

B.T.M. Holtus Managing Director J.M. McKenzie

Principal Certification Manager

First Issued: 17 August 2023 Date: 17 August 2023 Expiry date: 1 August 2028

© Integral publication of this certificate and adjoining reports is allowed

DEKRA Certification B.V. is Notified Body with ID no 0344

DEKRA Certification B.V. Meander 1051, 6825 MJ Arnhem P.O. Box 5185, 6802 ED Arnhem, The Netherlands T +31 88 96 83000 www.dekra.nl Company registration 09085396

Number: 6118114CE01

EU Quality Management System Certificate

Conformity Assessment Regulation 2017/745 on Medical devices, Annex IX Chapter I and III

This certificate covers the following device(s) / groups of device(s):

DIRECT DIGITAL X-RAY SYSTEMS (Z110311, class IIb)			
Intended Purpose: Mainly used for chest, abdomen, limbs and other parts (spine and pelvis) of X-ray photography examination, image for clinical diagnosis.			

Conditions for or limitations to the validity of this certificate:

N/A

Certificate History

Identification of the Common Specifications and Harmonized Standards complied with are documented within the technical documentation and audit assessments carried out. These are traceable through the DEKRA Certification B.V. Certification Notice. The Certification Notice also identifies the necessary information related to the quality management system of the manufacturer, including facilities.

Revision	Date of Issue certificate	Certification Notice Reference	///Action/
0	17 August 2023	////6096257CN03//////	/// /first/issue
	111111111111111111111111111111111111111		///////////////////////////////////////
		///////////////////////////////////////	///////////////////////////////////////

First Issued: 17 August 2023 Date: 17 August 2023 Expiry date: 1 August 2028

© Integral publication of this certificate and adjoining reports is allowed

DEKRA Certification B.V. is Notified Body with ID no 0344

DEKRA Certification B.V. Meander 1051, 6825 MJ Arnhem P.O. Box 5185, 6802 ED Arnhem, The Netherlands T +31 88 96 83000 www.dekra.nl Company registration 09085396