YIWEI Medical
Leader in AI Solutions for CNS

CE

EU DECLARATION OF CONFORMITY

Name and address of the manufacturer	YIWEI Medical Technology Co., Ltd Rooms 703-704, North Building, 3003 Shennan Middle Road, Futian District, Shenzhen, Guangdong, 518033, China
Name and address of the EU authorised representative Single Registration Number (SRN):	Crealkraft GmbH Burgfeldstr. 31, 45127 Essen DE-AR-000049943

We declare under our sole responsibility that

the medical device	**VR Cognitive Assessment and Rehab Training System** Baisc UDI-DI : 0697478241CognitiveZT *Applicable to any software version after V1* Model: Virtual Brain, Virtual Brain-A, Virtual Brain-R, Virtual Brain-H Virtual Brain-V, Virtual Brain-VA, Virtual Brain-VR, Virtual Brain-VH *Sterilization: n.a*
of class	**I** (Rule 11) according to annex VIII of the Regulation (EU) 2017/745

meets all requirements of the Medical Device Regulation (EU) 2017/745 that are applicable.

Applicable standards:
EN ISO 13485:2016 + A11:2021
EN ISO 14971:2019 + A11:2021
EN ISO 20417: 2021
EN 62304:2006 + A1:2015
ISO/IEC 25051:2014

Conformity assessment procedure	acc. to Annex IV of MDR
Notified Body	n.a.

Eric Ke

Shenzhen, 20.03.2026
Place, date

Eric Ke, PRRC
Name and function

Technical documentation verified by Suzhou Care-Real Medical Technology Co. LTD.

Crealkraft GmbH

Konformitätserklärung - Declaration of Conformity - 符合性声明Konformitätserklärung - Declaration of Conformity - 符合性声明Konformitätserklärung - Declaration of Conformity - 符合性声明