

EC DECLARATION OF CONFORMITY

Doc: HK-400IIIQPCE-06 Rev.: A4 Date: 2021-01-19



EC DECLARATION OF CONFORMITY

Name and address of the manufacturer: Name and address of the European Representative: We declare under our sole responsibility that	Shenzhen Hawk Medical Instrument Co. Ltd. 1st Floor, Building C, Jianyetai Industrial Zone, No.11 Minhuan Road, Fukang Community, Longhua Street, Longhua District, Shenzhen, 518109, Guangdong, P.R.China WellKang Ltd The Black Church, St. Mary's Place, Dublin 7, D07 P4AX, Ireland
the medical device:	Syringe Pump Model: HK-400III GMDN Code: 13217
of class:	Rule 11(3.2/7), IIb

e provisions of the directive 03/42/EEC and its transpositions in national laws which apply to it. The

meets the provisions of the directive 93/42/EEC and its transpositions in national laws which apply to it. The declaration is valid in connection with the "final inspection report" of the device.

according to annex IX of directive 93/42/EEC

Conformity assessment procedure:

Directive 93/42/EEC Annex II, excluding Section 4

Registration No.:

HD 2183512-1

Notified Body:

TÜV Rheinland LGA Products GmbH Tillystraße 2 90431 Nürnberg Deutschland CE 0197

The expiry date of the above mentioned declaration of conformity is May 26, 2024.

<u>Shenzhen, 2021-01-19</u> Place, date

General Manager

Name and function