MS
CM014

Certificate of Registration

Fuji Seiko Co., Ltd.

This is to certify that the above organization's Medical Devices Quality Management System conforms to Requirements of the following standard within the scope described in attached Appendix, and is registered by the JMAQA REGISTRATION CENTER as the result of the assessment.

Applicable Standard : JIS Q 13485:2018(ISO 13485:2016)

Registration No. : JMAQA-M036

Registration Date : 10 February 2020

Registration Expiry Date : 9 February 2026

Registration Revised Date : 21 November 2022

JAPAN MANAGEMENT ASSOCIATION QA REGISTRATION CENTER

Senior Executive Management
Masami Nakamura

3-1-22 Shiba-Koen Minato-ku Tokyo 105-8522,Japan

This is valid to be used in conjunction with attached Appendix.

MS
CM014

Appendix to Certificate of Registration

Appendix No. : JMAQA-M036-1　　Registration No. : JMAQA-M036

Fuji Seiko Co., Ltd.

860-6, Yamamiya, Fujinomiya, Shizuoka, Japan

Registration Scope :

Design, Development and Production of Precision Stainless Steel Tubing and Cannula for Medical Equipment

<The following sites are included.>

HEADQUARTERS / FUJINOMIYA FACTORY:

860-6, Yamamiya, Fujinomiya, Shizuoka, Japan

YAITA FACTORY:

7-1, Kobushi-dai, Yaita, Tochigi, Japan

Registration Revised Date: 21 November 2022