

# EC CERTIFICATE

## Full Quality Assurance System

Certificate No.: 248064-2017-CE-KOR-NA-PS Rev. 6.0

Project No.: PRJC-551628-2016-MSL-KOR

Valid Until: 28 May 2023

This is to certify that the quality system of:

**FINEMEDIX CO., LTD.**

140-9, Yuram-ro, Dong-gu, Daegu, 41059, Republic of Korea

For design, production and final product inspection/testing of:

**Endoscopic electric devices and Endoscopic non-electric devices**

Has been assessed with respect to:

**The conformity assessment procedure described in Annex II  
excluding section 4 of Council Directive 93/42/EEC on Medical  
Devices, as amended**

and found to comply

Further details of the product(s) and conditions for certification are given overleaf.

Place and date:  
Høvik, 21 April 2021

For the issuing office:  
Notified Body 2460  
DNV Product Assurance AS



*Eugenie Winger Husebye*  
Eugenie Winger Husebye  
Technical Reviewer

Notice: The Certificate is subject to terms and conditions as set out in the Certification Agreement. Failure to comply may render this Certificate invalid.

NOTIFIED BODY 2460: DNV Product Assurance AS, Veritasveien 3, 1363 Høvik, Norway, Tel +47 67 57 88 00, [www.dnv.com](http://www.dnv.com)

ICP-4-5-11-MDD-f2, rev.0

## Jurisdiction

Application of Council Directive 93/42/EEC of 14 June 1993, adopted as “Forskrift om Medisinsk Utstyr” by the Norwegian Ministry of Health and Care Services.

## Certificate history:

Revision	Description	Issue Date
0.0	Original Certificate	28 May 2019
1.0	Scope Extension_model added	04 December 2018
2.0	Product Name changed	18 February 2019
3.0	Scope Extension_model added	02 December 2019
4.0	Editorial change	17 December 2019
5.0	Scope Extension_model added	30 January 2020
<b>6.0</b>	<b>Site relocation</b>	<b>21 April 2021</b>

## Products covered by this Certificate:

Product Description	Product Name	Class
Endoscopic electric devices	ClearCut Knife	IIb
	ClearGrasp Snare	
	ClearCoajet	
	FineTome	
	ClearHemograsper	
	Clear-Hemostat	
	Clear-CoaBite	
Endoscopic non-electric devices	Clear-Jet Injection Catheter	IIa
	Clear-Bite Biopsy Forceps	
	Clear-Retriever	
	ClearTip	
	ClearEndoclip	
	Fine-Grab Basket	
Endoscopic non-electric devices	ClearCap Distal Attachment	Is

The complete list of devices is filed with the Notified Body



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### Sites covered by this certificate

Site Name	Address
FINEMEDIX CO., LTD.	140-9, Yuram-ro, Dong-gu, Daegu, 41059, Republic of Korea

### EU Representative

Medical Device Safety Service GmbH, Schiffgraben 41, 30175 Hannover, Germany



## Terms and conditions

The certificate is subject to the following terms and conditions:

- Any producer (see 2001/95/EC for a precise definition) is liable for damage caused by a defect in his product(s), in accordance with directive 85/374/EEC, as amended, concerning liability of defective products.
- The certificate is only valid for the products and/or manufacturing premises listed above.
- The Manufacturer shall fulfil the obligations arising out of the quality system as approved and uphold it so that it remains adequate and efficient.
- The Manufacturer shall inform the Notified Body of any intended updating of the quality system and the Notified Body will assess the changes and decide if the certificate remains valid.
- Periodical audits will be held, in order to verify that the Manufacturer maintains and applies the quality system. the Notified Body reserves the right, on a spot basis or based on suspicion, to pay unannounced visits.

The following may render this Certificate invalid:

- Changes in the quality system affecting production.
- Periodical audits not held within the allowed time window.

## Conformity declaration and marking of product

When meeting with the terms and conditions above, the producer may draw up an EC declaration of conformity and legally affix the CE mark followed by the Notified Body identification number.

End of Certificate