



# EC DESIGN EXAMINATION CERTIFICATE

## 93/42/EEC Directive of Medical Devices Annex II, Section 4

With the expire of the certificate M.2019.106.11991 the validity of the certificate  
M.2019.106.11991-1 will also end.

Company Name : YURIM MEDICAL Co., Ltd.

Company Address : 43, Jeongjung 3-gil, Osong-eup, Heungdeok-gu, Cheongju-si,  
Chungcheongbuk-do, Republic of Korea

Related Directives and Annex : 93/42/EEC Medical Devices Directive - Annex II (Section 4)

Product : Sterile P(LA/CL)[Poly(L-Lactide-co- $\epsilon$ -Caprolactone)]  
Suture with Needle - Class III

GMDN : 47193

Product Types are attached.

Certificate Number : M.2019.106.11991-1

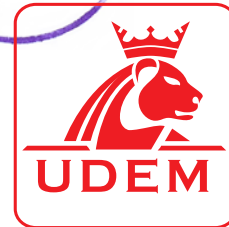
Report Number : MD.1013.IB

Initial Assessment Date : 13.09.2018

Registration Date : 31.05.2019

Revision Date /No : 25.05.2021/02

Expiry Date : 27.05.2024



The EC desing examination certificate refers to the above mentioned product. UDEM hereby declares that the requirements of Annex II, section 4 of the 93/42/EEC Directive have been met for the listed products. The above named manufacturer has established and applies a quality assurance system, which is subject to periodic surveillance audits, defined by Annex II, section 5 of the aforementioned directive. This certificate remains as the property of UDEM International Certification Auditing Training Centre Industry and Trade Inc. Co. to whom it must be returned upon request. The above named company and UDEM must keep a copy of this certificate for 5 years from the registration of the certificate. Usage of the CE mark is under the responsibility of the manufacturer with the completion of EC Declaration of Conformity. The above mentioned company must notify all changes related with the approved product to UDEM. If UDEM will not renew the expiry date of this certificate in question, the mentioned company should stop placing the product on the market. The validity of the certificate can be checked through [www.udem.com.tr](http://www.udem.com.tr).

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