



COMPANY WITH  
QUALITY SYSTEM  
CERTIFIED BY DNV  
ISO 13485

COMPANY WITH  
QUALITY SYSTEM  
CERTIFIED BY DNV  
ISO 9001

## BIO SUD MEDICAL SYSTEMS S.R.L. PRODOTTI E SISTEMI PER LA MEDICINA

S.r.l. Cap. Sociale € 52.000,00 i.v.  
C.F./P.IVA 03225090723 VAT IT03225090723  
Reg. Impr. N. 03225090723 N. iscrizione REA 249474 CCIAA di Bari

Via dei Fabbri, 23/25 – Zona Artig. ASI  
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### DECLARATION OF CONFORMITY

BIO SUD MEDICAL SYSTEMS S.r.l. headquartered in Modugno (BA), Via dei Fabbri 23/25, ITALY,  
MANUFACTURER of the Medical Device:

**COMMERCIAL NAME: BIOCRYL**

intended to be used as surgical absorbable suture

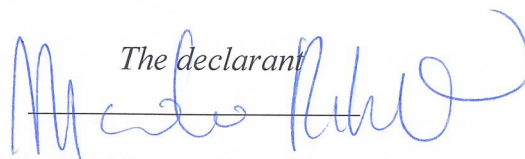
#### DECLARES

On its own responsibility, that the Medical device in question satisfies all the Essential Requirements mentioned in Annex I of 93/42/ECC Directive of Medical Devices and any subsequent amendments and additions.

For this purpose, it guarantees and declares on its own responsibility as follows:

- that the Medical Device in question satisfies the applicable provisions of 93/42/ECC Directive of Medical Devices and any subsequent amendments and additions, transposed into national legislation by Legislative Decree 24<sup>th</sup> February 1997, n. 46 and any subsequent amendments;
- that the Medical device in question is classified in Class III;
- that the Medical Device in question is provided STERILE;
- that the Medical Device is compliant with Essential Requirements and provisions of 93/42/EEC Directive of Medical Devices and any subsequent amendments and additions;
- that the Medical Device is manufactured according to the Quality System, that satisfies the Annex II's requirements of the above mentioned Legislative Decree, as indicated in the CE Certificate: Product Design Examination n. EPG-0165-18 (Expiry date 31/12/2027) and EC Declaration of Conformity Full Quality Assurance System n. QCT-0087-18 (Expiry date 31/12/2027), issued by "Istituto Superiore di sanità – Notified Body n.0373".

The manufacturer also declares to have established and maintain an appropriate procedure to guarantee the post-market surveillance, required by 93/42/ECC Directive and any subsequent amendments and additions.

  
*The declarant*  
(Nicola Ribatti – President of B.o.D.)  
**Bio Sud Medical Systems S.r.l.**