

BUREAU VERITAS
Certification



ID & CO S.r.l.

Via Lombardia, 10/d – 20098 San Giuliano Milanese (MI) - ITALY

Certified site:

Via Lombardia, 10/d – 20098 San Giuliano Milanese (MI) - ITALY

Bureau Veritas Italia S.p.A. certifies that the Production Quality Assurance System of the above organization has been audited and found to be in accordance with the requirements of

DIRECTIVE 93/42/EEC as amended

(according to Annex V)

In relation to the following products

Product subcategory :	Non active instruments
Generic group:	Accessories for operating room / Skin markers
Model:	PD01, PD01R, PD02, PD02R
Class:	Is

(may refer to the Annex of the certificate that lists all the products / models of devices subject to certification)

Reference BV practice: ZIG. N. 60649378

Original cycle start date: **10/04/2002**

Expiry date of previous cycle: **09/04/2020**

Certification / Recertification Audit date: **09/03/2020**

Certification / Recertification cycle start date: **09/04/2020**

Subject to the continued satisfactory operation of the organization's Management System, this certificate expires on: **26/05/2024**

Certificate No. - Version: IT273473 - 1

Revision date: **09/04/2020**


ANDREA FILIPPI – Certification SL Manager

This certificate is issued by Bureau Veritas Italia S.p.A. Viale Monza, 347-20126 Milan, as a notified body for the Directive 93/42/EEC, with identification number 1370

Further clarifications regarding the scope of this certificate and the applicability of the management system requirements may be obtained by consulting the organisation.
To check this certificate validity please refer to the website www.bureauveritas.it





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DIRECTIVE 93/42/EEC as amended

(according to Annex V)

In relation to the following products

Product subcategory :	Non active instruments
Generic group:	See annex
Model:	See annex
Class:	Is

(may refer to the Annex of the certificate that lists all the products / models of devices subject to certification)

Reference BV practice: ZIG. N. 60649378

Original cycle start date: **10/04/2002**

Expiry date of previous cycle: **09/04/2020**

Certification / Recertification Audit date: **09/03/2020**

Certification / Recertification cycle start date: **09/04/2020**

Subject to the continued satisfactory operation of the organization's Management System, this certificate expires on: **26/05/2024**

Certificate No. - Version: IT273472 - 1

Revision date: **09/04/2020**


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Annex to the CE Certificate
n° IT273472

Product subcategory :	Non active instruments
Generic group:	Accessories for operating room / Instruments sterile covers
Model:	CPxx/y, CPxx/x, CPxxx, CPxxx/x, CPxx/xxx, 61xx, 61xx y, 61 xx yy, 62xx, 2/6xxx, CPS xxx/x, CP y/y, CPMTxx, CPMTxxx, CP 75 IN T, CPS xxx, PMG 2xxx, PMG 4xxx, PMG 4xxx-xx, PMG 4xxx-xxx, PMGL 4xxx, PMGL 4xxx-xx, PMGL 4xxx-xxx, PMGXL 4xxx-xxx, PMGXL 4xxx-xx, PMG 5xxx, PMG 5xxx-xx, PMG 5xxx-xxx, PMGL 5xxx-xx, PMGL 5xxx-xxx, PMGXL 5xxx-xxx
Generic group:	Accessories for operating room / Instruments magnetic drapes
Model:	80710, 80720
Generic group:	Sterile abrasive tips cleaners
Model:	TC 1001
Class:	Is

Reference BV practice: ZIG. N. 60649378

Original cycle start date: **10/04/2002**

Expiry date of previous cycle: **09/04/2020**

Certification / Recertification Audit date: **09/03/2020**

Certification / Recertification cycle start date: **09/04/2020**

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Certificate No. - Version: IT273472 - 1

Revision date: **09/04/2020**


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Bureau Veritas Italia S.p.A. certifies that the Full Quality Assurance System of the above organization has been audited and found to be in accordance with the requirements of

DIRECTIVE 93/42/EEC as amended

(in accordance with Annex II - excluding paragraph 4)

In relation to the following products

Product subcategory :	Active surgical devices
Generic group:	See Annex
Model:	See Annex
Class:	IIb

(may refer to the Annex of the certificate that lists all the products / models of devices subject to certification)

Reference BV practice: ZIG. N. 60649378

Original cycle start date: **10/04/2002**

Expiry date of previous cycle: **09/04/2020**

Certification / Recertification Audit date: **09/03/2020**

Certification / Recertification cycle start date: **09/04/2020**

Subject to the continued satisfactory operation of the organization's Management System, this certificate expires on: **26/05/2024**

Certificate No. - Version: IT273477 - 1

Revision date: **09/04/2020**


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Annex to the CE Certificate
n° IT273477-1

Product subcategory :	Active surgical devices
Generic group:	Hand pieces for electrosurgical units
Model:	HCP-01, HCP-05, HCP-03, HCP-04.
Class:	IIb
Generic group:	Active electrodes (sterile and single use)
Model:	HCP-C0, HCP-C3, HCP-C4, HCP-C6, HCP-C7, HCP-C5, HCP-C8, HCP-C9, HCP-C1, HCP-C0T, HCP-C6T, HCP-C7T, HCP-C5T, HCP-F0, HCP-F3, HCP-F6, HCP-F1, HCP-F4, HCP-F8, HCP-F5.
Class:	IIb

Reference BV practice: ZIG. N. 60649378

Original cycle start date: **10/04/2002**
Expiry date of previous cycle: **09/04/2020**
Certification / Recertification Audit date: **09/03/2020**
Certification / Recertification cycle start date: **09/04/2020**

Subject to the continued satisfactory operation of the organization's Management System, this certificate expires on: **26/05/2024**

Certificate No. - Version: IT273477 - 1

Revision date: **09/04/2020**


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