

EC CERTIFICATE OF CONFORMITY 2026



This certificate of conformity conforms that the below mentioned product(s) and technical file as per Annex II has been examined, assessed and evaluated by competent appraisers of IFC according to declaration of conformity as per Annex II and found to be in conformance to the requirements of regulation **EU 2017/745** of the European parliament and of the Council of 5 April 2017 on medical devices, amending directive 2001/83/EC, regulation (EC) No 178/2002 and regulation (EC) No 1223/2009 and repealing council directives 90/385/EEC and 93/42/EEC. The details of Directive(s) and Standard mentioned below:

Issued To:

PROWA MEDICAL INSTRUMENTS

Manufacturer Address: P.O. Box 3036, 9-Km, Daska Road, Sialkot 51310-Pakistan

Brand Name: Prowa Medical Instruments

Product(s) Type: See Annex I

Regulation: Declaration of Conformity **EU 2017/745**

Class of MDR: Class 1

Harmonized European Standard References: ISO 14971:2019, ISO 13485:2016, ISO 9001:2015, ISO 15223:2021, BS EN 1041:2008+A1:2013, ISO 10993-1:2018, IEC 60601-1-Ed.3:2012.

Applied Directives: Regulation **EU 2017/745** of the European parliament and of the council of 5 April 2017 on medical devices.

Caution About CE Marking: The Label of the CE Marking should be not less then 5MM Height CE marking and EC Declaration of conformity are duties for the manufacturer or its applicant to puts the products on the market. He is responsible for initiating the CE marking and certification procedure as required by applicable law.

Note: If the devices is modified without the agreement of the undersigned, this declaration becomes invalid in relation to the modified product.

Certification is Based on the Technical File No: CE-EU-178

Certificate No: CE-EU-1014241

Latest Issue Date: January 12, 2026

Place and Date of Issue: Rome, January 12, 2026

Expiration Date: January 11, 2029

(This Certificate can be verified at <https://if-cb.com/validate-certificate/>)

Signed for and on the behalf of Italy branch:

Riccardo Basilio (Scheme Manager)



Head Office: 24 Helder St. South Croydon, CR2 6HT, United Kingdom

Italy Office: Via Dei Capocci, 39, 00184 Rome, RM, Italy

Web: www.if-cb.com | E-mail: italy@if-cb.com



EC CERTIFICATE OF CONFORMITY 2026



Company Name: PROWA MEDICAL INSTRUMENTS

Manufacturer Address: P.O. Box 3036, 9-Km, Daska Road, Sialkot 51310-Pakistan

Certificate Number: CE-EU-1014241

(Annex-I)

Serial No.	Product(s)Name	Serial No.	Product(s)Name	Serial No.	Product(s)Name
1	Wire Saws	28	Scissors	55	Nasal gouge
2	Uterine Tenaculum Forceps	29	Scalpels & Knives	56	Nasal files / Rasps
3	Uterine Scoops	30	Scalpels	57	Nasal Chisels
4	Uterine Curettes	31	Scalpel handles	58	Kidney Stone Forceps
5	Umbilical cord scissors	32	Root elevators	59	Intestinal grasping forceps
6	Tweezers	33	Rib Spreaders	60	Intestinal Clamps
7	Trocars	34	Rib Shears	61	Hooks
8	Tracheal Hooks	35	Rhinoplasty Knives	62	Hand Drills
9	Tooth extraction forceps	36	Retractors	63	Gall Stone forceps
10	Tonsil Snare	37	Retractors	64	Forceps
11	Tonsil Knives	38	Probes	65	Fixation Forceps
12	Tonsil Dissector	39	Periotomes	66	Endodontic explorers
13	Suture Forceps	40	Periosteal elevators	67	Elevators
14	Surgical Bulldog Clamps	41	Periodontal Curettes	68	Dissector
15	Spoons	42	Osteotomes	69	Dissecting forceps
16	Spatulas	43	Orthopaedic Rongeurs	70	Disimpaction Forceps
17	Skin hook	44	Needle Holders	71	Dentalfiles / rasp
18	Septum Elevators	45	Needle holders	72	Dental surgical scissors
19	Self-Retaining Retractor	46	Dental rongeurs	73	Dental scalers
20	Dental osteotomes	47	Chisels, Gouges	74	Bone Levers/elevator
21	Dental knives	48	Chin Retractor	75	Bone Holding Forceps
22	Dental excavators	49	Bone Spoons	76	Bone files / rasps
23	Curettes	50	Bone Saws	77	Bone Cutting Forceps
24	Bone Cutters	51	Bone Punches	78	Clamp manipulation forceps
25	Bone Curettes	52	ATS Instruments	79	Orthopaedic Instruments
26	Biopsy Forceps	53	Sterile Instruments	80	Sterilization Containers and Baskets
27	Abdominal Retractors	54	ENT Instruments		

This certificate remains the property of Inter Frontier Certification. Lack of full fulfillment of conditions as set out in the scheme rules may render this certificate invalid. The validity of the Inter Frontier Certification certificate will be maintained by annual surveillance audit and one renewal audit after three years. The use of accreditation mark indicates accreditation in respect of the activities covered by the scope of accreditation. This document is copyrighted and content may not be reproduced without Inter Frontier Certification prior written permission.



Signed for and on the behalf of Italy branch:

Riccardo Basilio (Scheme Manager)



Head Office: 24 Helder St. South Croydon, CR2 6HT, United Kingdom

Italy Office: Via Dei Capocci, 39, 00184 Rome, RM, Italy

Web: www.if-cb.com | E-mail: italy@if-cb.com

