



DECLARATION OF CONFORMITY

FULL QUALITY ASSURANCE SYSTEM

Product Name: Bipolar Forceps

Model Numbers: J1, J2, J3, J5, J7, J7D, J7U, J13, J14, J17, J18, J19, J20, J21, ACBF-018, ACBF-027

GMDN Code: Open-surgery electrosurgical handpiece/electrode, bipolar, reusable [47848]

Device Classification: IIb, Rule 9

Applied Standards:
EN 60601-2-2:2009 IEC 60601-2-2:2009
EN ISO 15223-1:2016 ISO 13485:2016
EN 1041:2008 EN ISO 14971:2012
ISO 10993-1:2009

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Notified Body & Identification Number: INTERTEK SEMKO AB
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EC Certificate Number: 41316578-04

Lot Number:

This declaration of conformity is issued under the sole responsibility of the manufacturer. The object of the declaration described above is in conformity with the relevant Community Harmonization Legislation. I hereby declare that the products specified above meet the requirements set forth in LVFS 2003:11 and Council Directive MDD 93/42/EEC, MDD Annex II (3) and conform to the identified harmonized standards. In addition, the object of the declaration described above is in conformity with RoHS Directive 2011/65/EU (RoHS 2) of the European Parliament and of the Council of 8 June 2011 on the restriction of the use of certain hazardous substances in electrical and electronic equipment. [RoHS]

Alan Ellman, CEO

February 12, 2025

Date