



**DECLARATION OF CONFORMITY**  
FULL QUALITY ASSURANCE SYSTEM

**Product Name:** Trigger-Flex

**Model Numbers:**

**GMDN Code:** Endoscopic electrosurgical handpiece/electrode, bipolar, single-use [57944]

**Device Classification:** IIb, Rule 9

**Applied Standards:** IEC 60601-1:2005/A1:2012      ISO 11135:2014  
 EN ISO 15223-1:2016      ISO 13485:2016  
 EN 1041:2008+A1:2013      EN ISO 14971:2012  
 IEC 60601-2-2:2009      ISO 10993-1:2009

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**Notified Body & Identification Number:** INTERTEK SEMKO AB  
 Kista, Sweden  
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**EC Certificate Number:** 41316578-03

**Lot #:**

*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

May 20, 2020  
 Date