



Declaration of Conformity

As Legal Manufacturer
We, 3M Company, Health Care Business,
3M Center, 2510 Conway Ave, Bldg. 275-5E-06
St. Paul, MN 55144 USA

hereby declare under our sole responsibility
that the CE marked products to which this declaration relates,

- 9661 3M™ Surgical Clipper with Pivot Head
- 9665 3M™ Charger, 240V (UK, Ireland)
- 9668 3M™ Charger, 230V (Europe)
- 9671 3M™ Surgical Clipper with Fixed Head
- 9675 3M™ Charger, 240V (UK, Ireland)
- 9678 3M™ Charger, 230V (Europe)
- 9677E Starter Kit (Europe)
- 9661L 3M™ Surgical Clipper with Pivot Head
- 9667L-E 3M™ Surgical Clipper Starter Kit (Includes 9661L and 9668L)
- 9665L 3M™ Drop-in charger stand with cord 100-240V (Plug Type G)
- 9668L 3M™ Drop-in charger stand with cord 100-240V (Plug Type C)

are classified,
per Rule 12 of Annex IX of the Medical Device Directive 93/42/EEC, as amended per 2007/47/EC,
as Class I active medical devices
and

are in accordance with Annex VII of Directive 93/42/EEC, as amended per 2007/47/EC,
on the approximation of the laws of the European Union Member States concerning medical devices

In addition, we declare that the above mentioned devices fulfill the applicable provisions of the Directive
93/42/EEC, as amended per 2007/47/EC
and

3M Health Care Business self-declares conformity with Directive 2011/65/EU 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 and compliance to the requirements of EN 50581:2012.

EU Representative Address
3M Deutschland GmbH
Health Care Business
Carl-Schurz-Str. 1
41453 Neuss, Germany

Signature:

Dianne Gibbs
3M Health Care
Division Regulatory Affairs Manager
Infection Prevention Division

Date:

18 July 2016