To whom it may concern

Ogni Dispositivo Medico, di Protezione e IVD deve essere registrato presso MHRA prima dell'immissione sul mercato in UK. Fabbricanti con sede fuori UK devono nominare un loro Rappresentante avente sede in UK (UKRP).

All medical devices, IVDs, PPE must be registered with the MHRA before they can be placed on the market in Great Britain. Non-UK manufacturer, are required to appoint a single UK Responsible Person (UKRP) within the UK.

La sottoscritta società/The undersigned company
THROAT SCOPE PTY LTD

con sede in/based in
LEVEL 25, 123 EAGLE ST., BRISBANE QLD 4000, AUSTRALIA

Dichiara/Declares

To have commissioned the company shown here to act as UKRP in his name and on behalf regarding the items listed below; that labels are updated within UK; that Letter of Designation that includes standard text provided by the MHRA was signed. All devices are registered with Public Access Database for Medical Device Registration.

ADVENA LTD, PURE OFFICES, PLATO CLOSE, WARWICK, CV34 6WE, UNITED KINGDOM

Elenco dei prodotti (Codice Gima + descrizione)
List of products (Gima code + description)

Codice Gima/Gima code	Nome/Name
	ThroatScope Handle x 1 (HH106)
	ThroatScope handle plus 50 depressors (TSH102)
	Tongue Depressor (pack of 50) (TS102)
	Pack of 10 Tongue Depressors

	(10PACK)
	TelScope single handle (TEL100)
	Telscope Shipper (48 single handles) (TEL100 X 48)

Date, place
24th May 2022, Cowbridge, UK

Stamp and signature