DECLARATION OF CONFORMITY

Regarding Medical Device Regulation (EU) 2017/745

Manufacturer:	Ningbo UNIMED Medical Instrument Co., Ltd.
Address:	26 Laoshan Road, Beilun, Ningbo, China.
EC Representative:	SUNGO Europe B.V.
Address:	Fascinatio Boulevard 522, Unit 1.7, 2909VA Capelle aan den IJssel, The Netherlands
Product Name:	Stethoscope
Model:	[SF100, SF101, SF207, SF208, SF402, SF500, SF601, SF701, SF702, SF200, SF201, SF202, SF203, SF205, SF306, SF411, SF412, SF413, SF501, SF502, SF503, SF504, SF507, SF508, SF512, SF513, SF602, SF603, SF5234, SF301, SF302, SF303, SF304, SF305.]
Classification:	Class I
Rule:	Rule 1, Annex VIII, Regulation (EU) 2017/745
Conformity Assessment Procedure:	Annex II+III of Regulation (EU) 2017/745
SRN:	CN-MF-000009819
Basic UDI-DI:	697454646STETHOSCOPEJV

We herewith declare that the above-mentioned products meet the requirements of Medical Device Regulation (EU) 2017/745 and the following harmonized standards.

EN ISO 14971:2019 EN ISO 15223-1:2021 EN ISO 20417:2021

Signature:

Name / Position: Zheng hui/ GM

Date: 2023.3.21

Place: Ningbo / China]

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