CERTIFICATE OF NOTIFICATION

This is to certify that, according to the council directive 93/42/EEC, SUNGO performed all notification duties and responsibilities as the European authorized representative of:

Applicant:
Address:

The Manufacturer has provided SUNGO with all the appropriate declarations according to the 93/42/EEC Directive requirements including the EC Declaration of Conformity confirming that his medical device, as stipulated here below, is fulfilling the applicable requirements of the European Council Directive 93/42/EEC.

Product(s):	Medical Face Mask
Type(s):	Non-sterile, ear loop, 17.5x9.5cm
Product Classification:	Class I

Where the manufacturer affix's the CE marking to the product listed they must ensure that all the requirements of the appropriate EU directive(s) have and continue to be met.

The notification of aforementioned device has been completed by the European Representative in United Kingdom. The UK Competent Authority is notified of the manufacturer's medical devices and has allocated registration. MHRA Registration number is CA017436

SUNGO
Executive Director
Sungo global service
CERTIFIED

Issued: Mar. 30 2020
Cert. No.: EU228518
Expiration Date: Mar. 29 2025

CE