Declaration of Conformity

Precision Medical Inc.
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EC REP

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CE 0088

REF

Precision Medical Air\Oxygen Low Flow Blenders;
PM5300EN, PM5300MEN, PM5300NIST, PM5300NISTAP, PM5300NISTAPMR

Precision Medical Air\Oxygen High Flow Blenders;
PM5200EN, PM5200MEN, PM5200NIST, PM5200NISTAP

Precision Medical Helium Oxygen, HeliO_2, 70\30% Low Flow Blenders;
PM5470EN, PM5470NIST

Precision Medical Helium Oxygen, HeliO_2, 80\20% Low Flow Blenders;
PM5480EN, PM5480NIST,

Precision Medical Helium Oxygen, HeliO_2, 70\30% High Flow Blenders;
PM5570EN, PM5570NIST

Precision Medical Helium Oxygen, HeliO_2, 80\20% High Flow Blenders;
PM5580EN, PM5580NIST

Classification: IIb

Classification criteria: Clause 3.2 Rule 11 of Annex IX of MDD

We hereby declare that an examination of the under mentioned production quality assurance system has been carried out following the requirements of the UK national legislation to which the undersigned is subjected, transposing Annex II, 3 of the Directive 93/42/EEC as amended by all subsequent directives.

Notified Body: Lloyd's Register LRQA
1 Trinity Park
Bickenhill Lane
Birmingham
B37 7ES
United Kingdom

Certification Registration No's: 10055119 Date of Expiry: 5 February 2023

Devices already manufactured: SN traceability via Device History Records
Validity of DOC: 6 February 2018 to Date of Expiry

Manufacture Representative: James Parker Signature: Digitally signed by James Parker Date: 2018.02.23 08:46:17 -05'00'

Position: Manager, Quality System/ISO Representative

Date of Issue: 6 February 2018