

EC Declaration of Conformity

Issued by

Medikro Oy

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Equipment Description:

This declaration of conformity is valid for the following product(s):

class IIa

M921	Medikro® SpiroStar DX (Serial) Spirometer
M921-E	Medikro® Serial OEM Spirometer
M921-WA, ref 703552	Spirometer kit (Serial)
M922	Medikro® SpiroMaster MX Spirometer
M929	Medikro® SpiroStar USB Spirometer
M929-E	Medikro® USB OEM Spirometer
M929-WA, ref 703554	Spirometer kit (USB)
M9831	Medikro® Spirometry Software (variants Smart, Pro and Sense)
M9228	SpiroSafe Disposable Flow Transducer
M9242	SpiroSafe Disposable Flow Transducer
M9242-WA, ref 703417	Disposable Flow Transducers (QTY 4)
M9242-WA, ref 703418	Disposable Flow Transducers (QTY 25)
M9242-WA, ref 703419	Disposable Flow Transducers (QTY 100)
M9472	Inhalation Trainer (Easyhaler PIF-meter)

class I accessories

M9227	Nose Clip
M9227-WA, ref 58550	Nose Clip
M9252	Pressure Tube for M9228 and SpiroStar
M9253	Pressure Tube for M9242 and SpiroStar
M9253-WA, ref 703415	Pressure Tube for M9242-WA and WA Spirometer
M9254	Disposable Tube for Easyhaler PIF-meter
M9255	Pressure Tube for M9242 and SpiroMaster MX
M9474	Calibration Syringe, volume 3000 ml
M9474-D	Calibration Syringe, volume 3000 ml, DYN'R OEM Model
M9477	Calibration Syringe, volume 1000 ml

(WA=Welch Allyn)

Applicable Directives:

The above products meet the essential requirements of the following directive(s):

Directive 93/42/EEC as amended by Directive 2007/47/EC

Place and Date
Kuopio, 18th February 2010



Mikko Eloranta
Managing Director

