EC DECLARATION OF CONFORMITY	Document Type:	Template
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According to:

LVFS 2001:7; Swedish legislation of EU Directive on In Vitro diagnostic Devices (98/79/EC) and amendments

1. **Type of equipment:**

Analyzing equipment for measuring Fractional Exhaled Nitric Oxide in exhaled breath in humans as a marker of inflammation

2. **Brand name or trade name:**

NIOX VERO

3. **Classification MDD/IVDD, class and rule:**

In Vitro Diagnostic Device 98/79/EC, other IVD

4. **Type designation(s)/Model no(s) and number of units:**

NIOX VERO, article number 12-1000, with regional configurations:

NIOX VERO (EU), article number 12-1100
NIOX VERO (US), article number 12-1200
NIOX VERO (JP), article number 12-1300
NIOX VERO (CN), article number 12-1400

NIOX VERO Breathing handle 12-1010

5. **Manufacturer's name, address, telephone and fax no:**

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As manufacturer we declare under sole responsibility that the equipment follows the provisions of the Directives stated above

Date and Place of issue

21-NOV-2016, Oxford

Name and signature of authorized person

Steve Harris, CEO