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:

Electrochemical sensor, replaceable accessory to NIOX VERO

2. Brand name or trade name:

NIOX VERO Sensor 60	12-1606
NIOX VERO Sensor 100	12-1610
NIOX VERO Sensor 300	12-1630
NIOX VERO Sensor 500	12-1650
NIOX VERO Sensor 1000	12-1700

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 sensors, M21

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

Circassia AB
Råsundavägen 18
SE-169 67 Solna
Sweden
tel +46 8 629 0780
fax +46 8 629 0781

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