VIAMED Ltd.
15 Station Road, Cross Hills, Keighley, West Yorkshire, BD20 7DT, UK.
Website: www.viamed.co.uk. Email: info@viamed.co.uk.
Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

VM-2160

hand held pulse oximeter for continuous and spot-check monitoring of functional arterial oxygen saturation (SpO2) and pulse rate,

Product No(s).

0012160

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
Cross Hills
Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

….....
Derek Lamb (Managing Director)

VIAMED Ltd.
15 Station Road, Cross Hills, Keighley, West Yorkshire, BD20 7DT, UK.
Website: www.viamed.co.uk. Email: info@viamed.co.uk.
Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

VM 2500-M and VM 2500-5

hand held capnograph and pulse oximeter for continuous and spot-check monitoring of end-tidal CO2 concentration (EtCO2), inspired CO2 concentration (FiCO2), functional arterial oxygen saturation (Sp02) respiration rate and pulse rate,

Product No(s).

4410500, 4410501, 4410520 and 4410521

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
Cross Hills
Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Derek Lamb (Managing Director)

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15 Station Road, Cross Hills, Keighley, West Yorkshire, BD20 7DT, UK.
Website: www.viamed.co.uk. Email: info@viamed.co.uk.
Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

VM-2160L

pulse oximeter for monitoring of functional artial oxygen satuation (SpO2) and pulse rate,

Product No(s).

0012155

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
Cross Hills
Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Derek Lamb (Managing Director)

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15 Station Road, Cross Hills, Keighley, West Yorkshire, BD20 7DT, UK.
Website: www.viamed.co.uk. Email: info@viamed.co.uk.
Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

VM-2105

Silicone Finger Oximeter for monitoring of functional arterial oxygen saturation (SpO2) and pulse rate,

Product No(s).

0012105 , 0012106

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
Cross Hills
Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Derek Lamb (Managing Director)

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Website: www.viamed.co.uk. Email: info@viamed.co.uk.
Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

4000 Series SpO2 Sensors

reusable and disposable pulse oximetry sensors for continuous and spot-check monitoring of functional arterial oxyen saturation (SpO2) and pulse rate.

Product No(s).

00149xx , 00150xx

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
Cross Hills
Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Derek Lamb (Managing Director)

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Apgar Timer

Timer used to gauge a given time interval when scoring newborn infants based on their vital signs and response to stimulus.

Product No(s).

0310100

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class I

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
Cross Hills
Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Derek Lamb (Managing Director)

VIAMED Ltd.
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Website: www.viamed.co.uk. Email: info@viamed.co.uk.
Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Cot Lid

Acrylic Sheet, hinged with or without louvers, to be used in conjunction with a basinet, to retain heat and humidity.

Product No(s).

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class I

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
Cross Hills
Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Derek Lamb (Managing Director)

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Website: www.viamed.co.uk. Email: info@viamed.co.uk.
Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Oxygen Sensor R-23v

A medical device for monitoring oxygen content in anaesthetic and intensive respiration devices and in incubators compatible to Drager 6850645 and 6809777.

Product No(s).

0110023

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIa

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
Cross Hills
Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Derek Lamb (Managing Director)

VIAMED Ltd.
15 Station Road, Cross Hills, Keighley, West Yorkshire, BD20 7DT, UK.
Website: www.viamed.co.uk. Email: info@viamed.co.uk.
Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Head Boxes

Enclosures to enable oxygen enriched air and humidity to be supplied to the patients head only.

Product No(s).

PP86xx

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIa

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
Cross Hills
Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

….....
Derek Lamb (Managing Director)

VIAMED Ltd.
15 Station Road, Cross Hills, Keighley, West Yorkshire, BD20 7DT, UK.
Website: www.viamed.co.uk. Email: info@viamed.co.uk.
Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Phototherapy Light Shield

Amber perspex head cover enabling photo-therapy to be administered whiles the eyes are protected.

Product No(s).

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIa

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
Cross Hills
Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Derek Lamb (Managing Director)

VIAMED Ltd.
15 Station Road, Cross Hills, Keighley, West Yorkshire, BD20 7DT, UK.
Website: www.viamed.co.uk. Email: info@viamed.co.uk.
Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Microstim DB III

Supramaximal Nerve Stimulator

Product No(s).

2510000

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIa

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
Cross Hills
Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Derek Lamb (Managing Director)

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable Tenthouses - Oxygen Hoods

Enclosures to enable oxygen enriched air and humidity to be supplied to the patients head only.

Product No(s).

23100xx

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIa

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
Cross Hills
Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Derek Lamb (Managing Director)

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15 Station Road, Cross Hills, Keighley, West Yorkshire, BD20 7DT, UK.
Website: www.viamed.co.uk. Email: info@viamed.co.uk.
Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Tom Thumb Infant Resuscitation

Hand Held, gas powered resuscitation unit.

Product No(s).

0310030/1/2/3 and 03100034, 0310072/4/5/6/ 0310080/2 0310092

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
Cross Hills
Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Derek Lamb (Managing Director)

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15 Station Road, Cross Hills, Keighley, West Yorkshire, BD20 7DT, UK.
Website: www.viamed.co.uk. Email: info@viamed.co.uk.
Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Finger Sensor - Adult. SF4001 NELLCOR Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two
wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014601

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
Cross Hills
Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

….....
Derek Lamb (Managing Director)

VIAMED Ltd.
15 Station Road, Cross Hills, Keighley, West Yorkshire, BD20 7DT, UK.
Website: www.viamed.co.uk. Email: info@viamed.co.uk.
Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Finger Sensor - Adult. SF4002 NELLCOR Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014602

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
Cross Hills
Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

….....
Derek Lamb (Managing Director)

VIAMED Ltd.
15 Station Road, Cross Hills, Keighley, West Yorkshire, BD20 7DT, UK.
Website: www.viamed.co.uk. Email: info@viamed.co.uk.
Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Finger Sensor - Adult. SF4006 HP/PHILIPS Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014603

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
Cross Hills
Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

….....
Derek Lamb (Managing Director)

VIAMED Ltd.
15 Station Road, Cross Hills, Keighley, West Yorkshire, BD20 7DT, UK.
Website: www.viamed.co.uk. Email: info@viamed.co.uk.
Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Finger Sensor - Adult. SF4007 HP/PHILIPS Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014604

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
Cross Hills
Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Derek Lamb (Managing Director)

VIAMED Ltd.
15 Station Road, Cross Hills, Keighley, West Yorkshire, BD20 7DT, UK.
Website: www.viamed.co.uk. Email: info@viamed.co.uk.
Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Finger Sensor - Adult. SF4011 DATEX Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014605

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
Cross Hills
Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Derek Lamb (Managing Director)

VIAMED Ltd.
15 Station Road, Cross Hills, Keighley, West Yorkshire, BD20 7DT, UK.
Website: www.viamed.co.uk. Email: info@viamed.co.uk.
Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Finger Sensor - Adult. SF4012 DATEX Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014606

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
Cross Hills
Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Derek Lamb (Managing Director)

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15 Station Road, Cross Hills, Keighley, West Yorkshire, BD20 7DT, UK.
Website: www.viamed.co.uk. Email: info@viamed.co.uk.
Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Finger Sensor - Adult. SF4016 OHMEDA Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014607

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
Cross Hills
Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Derek Lamb (Managing Director)

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15 Station Road, Cross Hills, Keighley, West Yorkshire, BD20 7DT, UK.
Website: www.viamed.co.uk. Email: info@viamed.co.uk.
Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Finger Sensor - Adult. SF4017 OHMEDA Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014608

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
Cross Hills
Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Derek Lamb (Managing Director)

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15 Station Road, Cross Hills, Keighley, West Yorkshire, BD20 7DT, UK.
Website: www.viamed.co.uk. Email: info@viamed.co.uk.
Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Finger Sensor - Adult. SF4021 BCI Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014609

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
Cross Hills
Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

….....
Derek Lamb (Managing Director)

VIAMED Ltd.
15 Station Road, Cross Hills, Keighley, West Yorkshire, BD20 7DT, UK.
Website: www.viamed.co.uk. Email: info@viamed.co.uk.
Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Finger Sensor - Adult. SF4026 NONIN Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014610

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
Cross Hills
Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Derek Lamb (Managing Director)

VIAMED Ltd.
15 Station Road, Cross Hills, Keighley, West Yorkshire, BD20 7DT, UK.
Website: www.viamed.co.uk. Email: info@viamed.co.uk.
Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Finger Sensor - Adult. SF4036 CSI Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014611

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
Cross Hills
Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Derek Lamb (Managing Director)

VIAMED Ltd.
15 Station Road, Cross Hills, Keighley, West Yorkshire, BD20 7DT, UK.
Website: www.viamed.co.uk. Email: info@viamed.co.uk.
Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Finger Sensor - Adult. SF4037 CSI Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014612

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
Cross Hills
Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

….....
Derek Lamb (Managing Director)

VIAMED Ltd.
15 Station Road, Cross Hills, Keighley, West Yorkshire, BD20 7DT, UK.
Website: www.viamed.co.uk. Email: info@viamed.co.uk.
Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Finger Sensor - Adult. SF4041 NOVAMETRIX Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014613

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
Cross Hills
Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

….....
Derek Lamb (Managing Director)

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Website: www.viamed.co.uk. Email: info@viamed.co.uk.
Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Finger Sensor - Adult. SF4046 NIHON KOHDEN Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014614

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Finger Sensor - Adult. SF4047 NIHON KOHDEN Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014615

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
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Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Website: www.viamed.co.uk. Email: info@viamed.co.uk.
Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Finger Sensor - Adult. SF4076 PALCO Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014621

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

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Issuer:

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Finger Sensor - Adult. SF4013 GE DATEX Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014625

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

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Keighley
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Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Pulse oximetry finger sensor SF 6500 VM VM 2160 Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014650

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

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15 Station Road
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Keighley
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United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Website: www.viamed.co.uk. Email: info@viamed.co.uk.
Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

CONSIGNED STOCK: SpO2 Finger Sensor 0014602 NELLCOR Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014692

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
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Keighley
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United Kingdom

Place, Date: Keighley , 5 Nov 2010.

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Silicone Finger Sensor - Adult. SC4001 NELLCOR 0.9M Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014701

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
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Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Silicone Finger Sensor - Adult. SC4002 NELLCOR 2M CABLE Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014702

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
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Keighley
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United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Silicone Finger Sensor - Adult. SC4006 HP/PHILIPS Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014703

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

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United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Silicone Finger Sensor - Adult. SC4007 HP/PHILIPS Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014704

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
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Keighley
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United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Silicone Finger Sensor - Adult. SC4011 DATEX Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014705

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

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Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Silicone Finger Sensor - Adult. SC4012 DATEX Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014706

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
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Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Website: www.viamed.co.uk. Email: info@viamed.co.uk.
Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Silicone Finger Sensor - Adult. SC4016 OHMEDA Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014707

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
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Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Silicone Finger Sensor - Adult. SC4017 OHMEDA Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014708

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

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United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Silicone Finger Sensor - Adult. SC4021 BCI Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014709

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

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Keighley
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United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Silicone Finger Sensor - Adult. SC4026 NONIN Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014710

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
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Class IIb

Application of the CE-Marking:

CE0086

Issuer:

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United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Silicone Finger Sensor - Adult. SC4036 CSI Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014711

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

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Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Silicone Finger Sensor - Adult. SC4037 CSI Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014712

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
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Class IIb

Application of the CE-Marking:

CE0086

Issuer:

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Keighley
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United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Silicone Finger Sensor - Adult. SC4041 NOVAMETRIX Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014713

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Silicone Finger Sensor - Adult. SC4046 NIHON KOHDEN Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014714

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Silicone Finger Sensor - Adult. SC4047 NIHON KOHDEN Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014715

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
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Class IIb

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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Silicone Finger Sensor - Adult. SC4076 PALCO Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014721

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
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Class IIb

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Silicone Finger Sensor - Adult. SC4031 MINOLTA 0.33M Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014722

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
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Class IIb

Application of the CE-Marking:

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United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

….....
Derek Lamb (Managing Director)

VIAMED Ltd.
15 Station Road, Cross Hills, Keighley, West Yorkshire, BD20 7DT, UK.
Website: www.viamed.co.uk. Email: info@viamed.co.uk.
Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Silicone Finger Sensor - Adult. HP/PHILIPS Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014723

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
Cross Hills
Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Derek Lamb (Managing Director)

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Silicone Finger Sensor - Adult. NIHON KOHDEN Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014724

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

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Issuer:

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Place, Date: Keighley , 5 Nov 2010.

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Silicone Finger Sensor - Adult. SC4013 GE DATEX Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014725

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
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Keighley
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United Kingdom

Place, Date: Keighley , 5 Nov 2010.

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Silicone Finger Sensor - Adult. SC4001VMP SAADAT 0.9M Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014740

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

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Issuer:

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Silicone Finger Sensor - Adult. SC6500VM VM 2160 Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014750

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

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Keighley
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Place, Date: Keighley , 5 Nov 2010.

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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Silicone Finger Sensor - Paediatric SCP6500VM VM 2160 Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014751

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
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Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Silicone Finger Sensor - Paediatric SCP4001 NELLCOR 0.9M Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014761

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
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Keighley
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United Kingdom

Place, Date: Keighley , 5 Nov 2010.

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Silicone Finger Sensor - Paediatric SCP4002 NELLCOR 2M CABLE Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014762

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

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Issuer:

Viamed Ltd.
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Keighley
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United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Silicone Finger Sensor - Paediatric SCP4006 HP/PHILIPS Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014763

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

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Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
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Keighley
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United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Silicone Finger Sensor - Paediatric SCP4007 HP/PHILIPS Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014764

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Silicone Finger Sensor - Paediatric SCP4011 DATEX Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014765

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
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Class IIb

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Silicone Finger Sensor - Paediatric SCP4012 DATEX Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014766

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
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Class IIb

Application of the CE-Marking:

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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Silicone Finger Sensor - Paediatric SCP4016 OHMEDA Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014767

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
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Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
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United Kingdom

Place, Date: Keighley , 5 Nov 2010.

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Silicone Finger Sensor - Paediatric SCP4017 OHMEDA Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014768

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

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Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Silicone Finger Sensor - Paediatric SCP4021 BCI Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014769

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

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Place, Date: Keighley , 5 Nov 2010.

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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Silicone Finger Sensor - Paediatric SCP4026 NONIN Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014770

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
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CE0086

Issuer:

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United Kingdom

Place, Date: Keighley , 5 Nov 2010.

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Silicone Finger Sensor - Paediatric SCP4036 CSI Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014771

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Silicone Finger Sensor - Paediatric SCP4037 CSI Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014772

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

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CE0086

Issuer:

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Keighley
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United Kingdom

Place, Date: Keighley , 5 Nov 2010.

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Silicone Finger Sensor - Paediatric SCP4041 NOVAMETRIX Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014773

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
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Class IIb

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CE0086

Issuer:

Viamed Ltd.
15 Station Road
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Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Silicone Finger Sensor - Paediatric SCP4046 NIHON KOHDEN Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014774

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
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Class IIb

Application of the CE-Marking:

CE0086

Issuer:

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United Kingdom

Place, Date: Keighley , 5 Nov 2010.

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Silicone Finger Sensor - Paediatric SCP4047 NIHON KOHDEN Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014775

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

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CE0086

Issuer:

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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Silicone Finger Sensor - Paediatric SCP4061 DATASCOPE 0.9M Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014777

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

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Place, Date: Keighley , 5 Nov 2010.

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Silicone Finger Sensor - Paediatric SCP4062 DATASCOPE 2.95M Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014778

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
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Class IIb

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United Kingdom

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Silicone Finger Sensor - Paediatric SCP4031 MINOLTA 0.33M Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014782

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

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CE0086

Issuer:

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15 Station Road
Cross Hills
Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

….....
Derek Lamb (Managing Director)

VIAMED Ltd.
15 Station Road, Cross Hills, Keighley, West Yorkshire, BD20 7DT, UK.
Website: www.viamed.co.uk. Email: info@viamed.co.uk.
Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Silicone Finger Sensor - Paediatric SCP4001 SAADAT 0.9M Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014790

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
Cross Hills
Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Derek Lamb (Managing Director)

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Website: www.viamed.co.uk. Email: info@viamed.co.uk.
Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Pulse oximetry Ear Sensor EP4001 NELLCOR 0.9 Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014801

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
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Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

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Derek Lamb (Managing Director)

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Website: www.viamed.co.uk. Email: info@viamed.co.uk.
Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Pulse oximetry Ear Sensor EP4007 HP/PHILIPS 2.0 Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014802

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
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Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Derek Lamb (Managing Director)

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15 Station Road, Cross Hills, Keighley, West Yorkshire, BD20 7DT, UK.
Website: www.viamed.co.uk. Email: info@viamed.co.uk.
Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Pulse oximetry Ear Sensor EP4012 DATEX 2.95M Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014804

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
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Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Derek Lamb (Managing Director)

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Website: www.viamed.co.uk. Email: info@viamed.co.uk.
Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Pulse oximetry Ear Sensor EP4016 OHMEDA 2.95M Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014805

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

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Keighley
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Place, Date: Keighley , 5 Nov 2010.

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Pulse Oximetry Wrap Sensor W4001 SAADAT 0.9M Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014820

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

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Issuer:

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Pulse Oximetry Wrap Sensor W4001 NELLCOR 0.9M Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014821

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

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Keighley
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United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Website: www.viamed.co.uk. Email: info@viamed.co.uk.
Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Pulse Oximetry Wrap Sensor W4002 NELLCOR 2.0M Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014822

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

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Keighley
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United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Pulse Oximetry Wrap Sensor W4006 HP/PHILIPS 2.95M Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014823

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

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Issuer:

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United Kingdom

Place, Date: Keighley , 5 Nov 2010.

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Pulse Oximetry Wrap Sensor W4007 HP/PHILIPS 2.0M Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014824

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

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Issuer:

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Place, Date: Keighley , 5 Nov 2010.

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Pulse oximetry Ear Sensor EP6500VM VM 2160 Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014825

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
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Class IIb

Application of the CE-Marking:

CE0086

Issuer:

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Place, Date: Keighley , 5 Nov 2010.

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Pulse Oximetry Wrap Sensor W4011 DATEX 0.90M Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014826

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
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Class IIb

Application of the CE-Marking:

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Pulse Oximetry Wrap Sensor W4012 DATEX 2.95M Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014827

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

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15 Station Road
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Keighley
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United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Website: www.viamed.co.uk. Email: info@viamed.co.uk.
Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Pulse Oximetry Wrap Sensor W4016 OHMEDA 2.95M Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014828

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

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15 Station Road
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Keighley
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United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Derek Lamb (Managing Director)

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Pulse Oximetry Wrap Sensor W4017 OHMEDA 1.00M Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014829

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

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United Kingdom

Place, Date: Keighley , 5 Nov 2010.

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Pulse Oximetry Wrap Sensor W6500VM VM 2160 Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014835

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
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Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Derek Lamb (Managing Director)

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15 Station Road, Cross Hills, Keighley, West Yorkshire, BD20 7DT, UK.
Website: www.viamed.co.uk. Email: info@viamed.co.uk.
Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Pulse oximetry multi-site Y sensor VM 2160 Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014845

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

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15 Station Road
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Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Derek Lamb (Managing Director)

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Pulse oximetry extension cable 1.2 m XT 6500 VM 2160 Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014895

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
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Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Derek Lamb (Managing Director)

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Pulse oximetry extension cable 2.4 m XT 6501 VM 2160 Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014896

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
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Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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15 Station Road, Cross Hills, Keighley, West Yorkshire, BD20 7DT, UK.
Website: www.viamed.co.uk. Email: info@viamed.co.uk.
Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Sample pack. 4000 SAMPLE PACK Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014900

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

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15 Station Road
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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Adult. 4000 MICRO FOAM 1-AF NELLCOR Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014901

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Paediatric. 4000 MICRO FOAM 1-PF NELLCOR Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014902

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

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15 Station Road
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Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Infant. 4000 MICRO FOAM 1-IF NELLCOR Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014903

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
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Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Neonatal. 4000 MICRO FOAM 1-NF NELLCOR Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014904

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

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Cross Hills
Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

….....
Derek Lamb (Managing Director)

VIAMED Ltd.
15 Station Road, Cross Hills, Keighley, West Yorkshire, BD20 7DT, UK.
Website: www.viamed.co.uk. Email: info@viamed.co.uk.
Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Adult. 4000 TRANSPORE 1-AT NELLCOR Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014905

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
Cross Hills
Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Derek Lamb (Managing Director)

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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Paediatric. 4000 TRANSPORE 1-PT NELLCOR Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014906

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
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Keighley
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Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

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Derek Lamb (Managing Director)

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Adult. 4000 PLASTER 1-AP NELLCOR Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014909

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
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Keighley
West Yorkshire, BD20 7DT
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Place, Date: Keighley , 5 Nov 2010.

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Derek Lamb (Managing Director)

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Paediatric. 4000 PLASTER 1-PP NELLCOR Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014910

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

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Place, Date: Keighley , 5 Nov 2010.

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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Infant. 4000 PLASTER 1-IP NELLCOR Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014911

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

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Issuer:

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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Neonatal. 4000 PLASTER 1-NP NELLCOR Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014912

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

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Application of the CE-Marking:

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Issuer:

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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Adult. 4000 MICRO FOAM 2-AF OHMEDA Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014913

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

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Application of the CE-Marking:

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Issuer:

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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Paediatric. 4000 MICRO FOAM 2-PF OHMEDA Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014914

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

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Application of the CE-Marking:

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Issuer:

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Place, Date: Keighley , 5 Nov 2010.

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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Infant. 4000 MICRO FOAM 2-IF OHMEDA Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014915

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

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Issuer:

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Place, Date: Keighley , 5 Nov 2010.

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Neonatal. 4000 MICRO FOAM 2-NF OHMEDA Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014916

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Adult. 4000 TRANSPORE 2-AT OHMEDA Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014917

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Paediatric. 4000 TRANSPORE 2-PT OHMEDA Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014918

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Adult. 4000 PLASTER 2-AP OHMEDA Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014921

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

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Issuer:

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Paediatric. 4000 PLASTER 2-PP OHMEDA Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014922

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Infant. 4000 PLASTER 2-IP OHMEDA Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014923

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Neonatal. 4000 PLASTER 2-NP OHMEDA Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014924

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Adult. 4000 MICRO FOAM 3-AF DATEX Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014925

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

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Issuer:

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Paediatric. 4000 MICRO FOAM 3-PF DATEX Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014926

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

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Issuer:

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Infant. 4000 MICRO FOAM 3-IF DATEX Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014927

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Neonatal. 4000 MICRO FOAM 3-NF DATEX Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014928

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Adult. 4000 TRANSPORE 3-AT DATEX Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014929

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Paediatric. 4000 TRANSPORE 3-PT DATEX Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014930

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Adult. 4000 PLASTER 3-AP DATEX Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014933

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Paediatric. 4000 PLASTER 3-PP DATEX Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014934

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
Cross Hills
Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

….....
Derek Lamb (Managing Director)

VIAMED Ltd.
15 Station Road, Cross Hills, Keighley, West Yorkshire, BD20 7DT, UK.
Website: www.viamed.co.uk. Email: info@viamed.co.uk.
Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Infant. 4000 PLASTER 3-IP DATEX Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014935

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Neonatal. 4000 PLASTER 3-NP DATEX Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014936

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

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Issuer:

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Place, Date: Keighley , 5 Nov 2010.

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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Adult. 4000 MICRO FOAM 4-AF CSI Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014937

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

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Application of the CE-Marking:

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Issuer:

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Place, Date: Keighley , 5 Nov 2010.

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Paediatric. 4000 MICRO FOAM 4-PF CSI Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014938

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Infant. 4000 MICRO FOAM 4-IF CSI Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014939

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Neonatal. 4000 MICRO FOAM 4-NF CSI Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014940

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Adult. 4000 TRANSPORE 4-AT CSI Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014941

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Paediatric. 4000 TRANSPORE 4-PT CSI Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014942

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

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Place, Date: Keighley , 5 Nov 2010.

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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Adult. 4000 PLASTER 4-AP CSI Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014945

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Paediatric. 4000 PLASTER 4-PP CSI Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014946

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
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Class IIb

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Issuer:

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Place, Date: Keighley , 5 Nov 2010.

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Infant. 4000 PLASTER 4-IP CSI Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014947

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

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Issuer:

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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Neonatal. 4000 PLASTER 4-NP CSI Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014948

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
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Class IIb

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Issuer:

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Adult. 4000 MICRO FOAM 5-AF BCI Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014949

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

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Keighley
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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Paediatric. 4000 MICRO FOAM 5-PF BCI Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014950

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Infant. 4000 MICRO FOAM 5-IF BCI Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014951

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Neonatal. 4000 MICRO FOAM 5-NF BCI Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014952

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Adult. 4000 TRANSPORE 5-AT BCI Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014953

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Paediatric. 4000 TRANSPORE 5-PT BCI Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014954

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Adult. 4000 PLASTER 5-AP BCI Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014957

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Paediatric. 4000 PLASTER 5-PP BCI Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014958

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Infant. 4000 PLASTER 5-IP BCI Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014959

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Neonatal. 4000 PLASTER 5-NP BCI Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014960

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

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Application of the CE-Marking:

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Adult. 4000 MICRO FOAM 6-AF NOVAMETRIX Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014961

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Paediatric. 4000 MICRO FOAM 6-PF NOVAMETRIX Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014962

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
Cross Hills
Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

….....
Derek Lamb (Managing Director)

VIAMED Ltd.
15 Station Road, Cross Hills, Keighley, West Yorkshire, BD20 7DT, UK.
Website: www.viamed.co.uk. Email: info@viamed.co.uk.
Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Infant. 4000 MICRO FOAM 6-IF NOVAMETRIX Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014963

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Neonatal. 4000 MICRO FOAM 6-NF NOVAMETRIX Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014964

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Adult. 4000 TRANSPORE 6-AT NOVAMETRIX Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014965

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Paediatric. 4000 TRANSPORE 6-PT NOVAMETRIX Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014966

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Adult. 4000 PLASTER 6-AP NOVAMETRIX Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014969

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Paediatric. 4000 PLASTER 6-PP NOVAMETRIX Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014970

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Infant. 4000 PLASTER 6-IP NOVAMETRIX Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014971

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Neonatal. 4000 PLASTER 6-NP NOVAMETRIX Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014972

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Adult. 4000 MICRO FOAM 7-AF NONIN Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014973

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Paediatric. 4000 MICRO FOAM 7-PF NONIN Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014974

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Infant. 4000 MICRO FOAM 7-IF NONIN Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014975

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Neonatal. 4000 MICRO FOAM 7-NF NONIN Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014976

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

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Place, Date: Keighley , 5 Nov 2010.

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Adult. 4000 TRANSPORE 7-AT NONIN Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014977

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

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Issuer:

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Paediatric. 4000 TRANSPORE 7-PT NONIN Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014978

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Adult. 4000 PLASTER 7-AP NONIN Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014981

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Paediatric. 4000 PLASTER 7-PP NONIN Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014982

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Infant. 4000 PLASTER 7-IP NONIN Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014983

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Neonatal. 4000 PLASTER 7-NP NONIN Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014984

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Adult. 4000 MICRO FOAM 8-AF VIAMED Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014985

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Paediatric. 4000 MICRO FOAM 8-PF VIAMED Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014986

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Infant. 4000 MICRO FOAM 8-IF VIAMED Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014987

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Neonatal. 4000 MICRO FOAM 8-NF VIAMED Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014988

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Adult. 4000 TRANSPORE 8-AT VIAMED Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014989

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
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Derek Lamb (Managing Director)

VIAMED Ltd.
15 Station Road, Cross Hills, Keighley, West Yorkshire, BD20 7DT, UK.
Website: www.viamed.co.uk. Email: info@viamed.co.uk.
Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Paediatric. 4000 TRANSPORE 8-PT VIAMED Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014990

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.
15 Station Road
Cross Hills
Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date: Keighley , 5 Nov 2010.

Legally binding signature:

….....
Derek Lamb (Managing Director)

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Website: www.viamed.co.uk. Email: info@viamed.co.uk.
Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Adult. 4000 PLASTER 8-AP VIAMED Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014993

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIb

Application of the CE-Marking:

CE0086

Issuer:

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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Paediatric. 4000 PLASTER 8-PP VIAMED Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014994

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Infant. 4000 PLASTER 8-IP VIAMED Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014995

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

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Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582.

EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Disposable SpO2 Sensors - Neonatal. 4000 PLASTER 8-NP VIAMED Compatible

For use with pulse oximeters in oerder to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014996

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.
In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

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