[redacted]

SpO_2 probe - pulse oximetry sensor:

For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.

I the undersigned certify that the part(s) listed **P878RA**
is/are compatible with the manufacturer(s) part (s) **INVIVO®** **9383**

It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.

This declaration is supported by:

Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086

Signed: Date: 23 April 1999

Position: Managing Director

Viamed Ltd. 15 Station Road , Crosshills,
Keighley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Web www.viamed.co.uk

SpO_2 probe - pulse oximetry sensor:

For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.

I the undersigned certify that the part(s) listed **P886RA**
is/are compatible with the manufacturer(s) part (s) **KONTRON®** **0608010**

It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.

This declaration is supported by:

Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086

Signed: Date: 23 April 1999

Position: Managing Director

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Keighley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Web www.viamed.co.uk

[redacted]

SpO_2 probe - pulse oximetry sensor:

For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.

I the undersigned certify that the part(s) listed **P875RA**
is/are compatible with the manufacturer(s) part (s) **SPACELABS®** **369083-001**

It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.

This declaration is supported by:

Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
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Signed: Date: 23 April 1999

Position: Managing Director

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Keighley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Web www.viamed.co.uk

[redacted]

SpO_2 probe - pulse oximetry sensor:

For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.

I the undersigned certify that the part(s) listed **P875RA**
is/are compatible with the manufacturer(s) part (s) **Novametrics®** **8660**

It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.

This declaration is supported by:

Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
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Notified body number CE 0086

Signed: Date: 23 April 1999

Position: Managing Director

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Keighley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Web www.viamed.co.uk

[redacted]

SpO_2 probe - pulse oximetry sensor:

For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.

I the undersigned certify that the part(s) listed **P867YA**
is/are compatible with the manufacturer(s) part (s) **OHMEDA®** **Flexible probes**

It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.

This declaration is supported by:

Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
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Notified body number CE 0086

Signed: Date: 23 April 1999

Position: Managing Director

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Keighley, West Yorkshire, BD20 7DT. UK.
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Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086
Signed: Date: 23 April 1999
Position: Managing Director

Viamed Ltd. 15 Station Road , Crosshills,
Keigley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Webb www.viamed.co.uk cocP973e10

Certificate of Compatibility
Class IIb

SpO2 probe - pulse oximetry sensor:
For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.
I the undersigned certify that the part(s) listed P973E5
is/are compatible with the manufacturer(s) part (s) Datex® SAS-C1
It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.
This declaration is supported by:
Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086
Signed: Date: 23 April 1999
Position: Managing Director

Viamed Ltd. 15 Station Road , Crosshills,
Keigley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Webb www.viamed.co.uk cocP956E8

Certificate of Compatibility
Class IIb

SpO2 probe - pulse oximetry sensor:
For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.
I the undersigned certify that the part(s) listed P973E10
is/are compatible with the manufacturer(s) part (s) Datex® SAS-C3
It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.
This declaration is supported by:
Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086
Signed: Date: 23 April 1999
Position: Managing Director

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Keigley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Webb www.viamed.co.uk cocP973e10

Certificate of Compatibility
Class IIb

SpO2 probe - pulse oximetry sensor:
For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.
I the undersigned certify that the part(s) listed P973E5
is/are compatible with the manufacturer(s) part (s) Datex® SAS-C1
It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.
This declaration is supported by:
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EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086
Signed: Date: 23 April 1999
Position: Managing Director

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Keigley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Webb www.viamed.co.uk cocP973E5
Certificate of Compatibility
Class IIb

SpO2 probe - pulse oximetry sensor:
For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.
I the undersigned certify that the part(s) listed P973E10
is/are compatible with the manufacturer(s) part (s) Datex® SAS-C3
It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.
This declaration is supported by:
Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)

Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086
Signed: Date: 23 April 1999
Position: Managing Director

Viamed Ltd. 15 Station Road , Crosshills,
Keigley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Webb www.viamed.co.uk cocP956e4

Certificate of Compatibility
Class IIb

SpO2 probe - pulse oximetry sensor:
For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.
I the undersigned certify that the part(s) listed P956E4
is/are compatible with the manufacturer(s) part (s) Nellcor® EC4
It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.
This declaration is supported by:
Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086
Signed: Date: 23 April 1999
Position: Managing Director

Viamed Ltd. 15 Station Road , Crosshills,
Keigley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Webb www.viamed.co.uk cocP956E4

Certificate of Compatibility
Class IIb

SpO2 probe - pulse oximetry sensor:
For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.
I the undersigned certify that the part(s) listed P956E8
is/are compatible with the manufacturer(s) part (s) Nellcor® EC8
It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.
This declaration is supported by:
Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086
Signed: Date: 23 April 1999
Position: Managing Director

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Keigley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Webb www.viamed.co.uk cocP956e8

Certificate of Compatibility
Class IIb

SpO2 probe - pulse oximetry sensor:
For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.
I the undersigned certify that the part(s) listed P956E4
is/are compatible with the manufacturer(s) part (s) Nellcor® EC8
It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.
This declaration is supported by:
Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086
Signed: Date: 23 April 1999
Position: Managing Director

Viamed Ltd. 15 Station Road , Crosshills,
Keigley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Webb www.viamed.co.uk cocP956E8
Certificate of Compatibility
Class IIb

SpO2 probe - pulse oximetry sensor:
For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.
I the undersigned certify that the part(s) listed P956E8
is/are compatible with the manufacturer(s) part (s) Nellcor® EC8
It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.
This declaration is supported by:
Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)

Certificate of Compatibility
Class IIb

SpO2 probe - pulse oximetry sensor:

For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.

I the undersigned certify that the part(s) listed P871RA
is/are compatible with the manufacturer(s) part (s) NoninR 8604K, 8000K2, 8000AA-1

It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.

This declaration is supported by:

Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086

Signed: Date: 18 April 2001

Position: Managing Director

Viamed Ltd. 15 Station Road , Crosshills,
Keigley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Web www.viamed.co.uk

Certificate of Compatibility
Class IIb

SpO_2 probe - pulse oximetry sensor extension cable:

For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.

I the undersigned certify that the part(s) listed **P971E3**
is/are compatible with the manufacturer(s) part (s) **Nonin**

It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.

This declaration is supported by:

Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086

Signed: Date:25th November 2003

Position: Managing Director

VIAMED

Viamed Ltd. 15 Station Road , Crosshills,
Keigley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Web www.viamed.co.uk

SpO_2 probe - pulse oximetry sensor:

For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.

I the undersigned certify that the part(s) listed **P867RA**
is/are compatible with the manufacturer(s) part (s) **OHMEDA® Finger probes 380-1000-042**

It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.

This declaration is supported by:

Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086

Signed: Date: 23 April 1999

Position: Managing Director

Viamed Ltd. 15 Station Road , Crosshills,
Keighley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Web www.viamed.co.uk

Certificate of Compatibility
Class IIb

SpO_2 probe - pulse oximetry sensor extension cable:

For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.

I the undersigned certify that the part(s) listed **P991E10**
is/are compatible with the manufacturer(s) part (s) **Critikon**

It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.

This declaration is supported by:

Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086

Signed: Date:25th November 2003

Position: Managing Director

VIAMED

Viamed Ltd. 15 Station Road , Crosshills,
Keigley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Web www.viamed.co.uk

SpO_2 probe - pulse oximetry sensor:

For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.

I the undersigned certify that the part(s) listed **P861RA**
is/are compatible with the manufacturer(s) part (s) **BCI®** **3044**

It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.

This declaration is supported by:

Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086

Signed: Date: 23 April 1999

Position: Managing Director

Viamed Ltd. 15 Station Road , Crosshills,
Keighley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Web www.viamed.co.uk

Certificate of Compatibility
Class IIb

SpO_2 probe - pulse oximetry sensor extension cable:

For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.

I the undersigned certify that the part(s) listed **P971E8**
is/are compatible with the manufacturer(s) part (s) **Nonin**

It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.

This declaration is supported by:

Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086

Signed: Date:25th November 2003

Position: Managing Director

VIAMED

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Keigley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Web www.viamed.co.uk

Certificate of Compatibility
Class IIb

SpO_2 probe - pulse oximetry sensor extension cable:

For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.

I the undersigned certify that the part(s) listed **P969E8**
is/are compatible with the manufacturer(s) part (s) **Criticare**

It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.

This declaration is supported by:

Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086

Signed: Date:25th November 2003

Position: Managing Director

VIAMED

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email info@viamed.co.uk Web www.viamed.co.uk

Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086
Signed: Date: 23 April 1999
Position: Managing Director

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Keigley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Webb www.viamed.co.uk cocP973e10

Certificate of Compatibility
Class IIb

SpO2 probe - pulse oximetry sensor:
For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.
I the undersigned certify that the part(s) listed P973E10
is/are compatible with the manufacturer(s) part (s) Datex® SAS-C3
It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.
This declaration is supported by:
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EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086
Signed: Date: 23 April 1999
Position: Managing Director

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email info@viamed.co.uk Webb www.viamed.co.uk cocP956E10

Certificate of Compatibility
Class IIb

SpO2 probe - pulse oximetry sensor:
For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.
I the undersigned certify that the part(s) listed P956E4
is/are compatible with the manufacturer(s) part (s) Nellcor® EC4
It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.
This declaration is supported by:
Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086
Signed: Date: 23 April 1999
Position: Managing Director

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Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Webb www.viamed.co.uk cocP956e4

Certificate of Compatibility
Class IIb

SpO2 probe - pulse oximetry sensor:
For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.
I the undersigned certify that the part(s) listed P956E4
is/are compatible with the manufacturer(s) part (s) Nellcor® EC
It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.
This declaration is supported by:
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email info@viamed.co.uk Webb www.viamed.co.uk cocP956E4
Certificate of Compatibility
Class IIb

SpO2 probe - pulse oximetry sensor:
For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.
I the undersigned certify that the part(s) listed P956E4
is/are compatible with the manufacturer(s) part (s) Nellcor® EC4
It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.
This declaration is supported by:
Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)

Certificate of Compatibility
Class IIb

SpO_2 probe - pulse oximetry sensor:

For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.

I the undersigned certify that the part(s) listed **P863RA**
is/are compatible with the manufacturer(s) part (s) **Datascope[R]** **0600-00-0026-01**

It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.

This declaration is supported by:

Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086

Signed: Date: 23 April 1999

Position: Managing Director

VIAMED

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Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Web www.viamed.co.uk

Certificate of Compatibility
Class IIb

SpO_2 probe - pulse oximetry sensor extension cable:

For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.

I the undersigned certify that the part(s) listed **P973E10**
is/are compatible with the manufacturer(s) part (s) **Datex**

It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.

This declaration is supported by:

Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086

Signed: Date:25th November 2003

Position: Managing Director

VIAMED

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Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Web www.viamed.co.uk

Certificate of Compatibility
Class IIb

SpO_2 probe - pulse oximetry sensor extension cable:

For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.

I the undersigned certify that the part(s) listed **P968E8**
is/are compatible with the manufacturer(s) part (s) **Criticare**

It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.

This declaration is supported by:

Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086

Signed: Date:25th November 2003

Position: Managing Director

VIAMED

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Keigley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Web www.viamed.co.uk

Certificate of Compatibility
Class IIb

SpO_2 probe - pulse oximetry sensor extension cable:

For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.

I the undersigned certify that the part(s) listed **P963E10**
is/are compatible with the manufacturer(s) part (s) **Datascope**

It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.

This declaration is supported by:

Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086

Signed: Date:25th November 2003

Position: Managing Director

VIAMED

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Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Web www.viamed.co.uk

[redacted]

SpO_2 probe - pulse oximetry sensor:

For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.

I the undersigned certify that the part(s) listed **P860RA**
is/are compatible with the manufacturer(s) part (s) **Simed®/Baxter ®** **600333-003**

It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.

This declaration is supported by:

Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086

Signed: Date: 23 April 1999

Position: Managing Director

Viamed Ltd. 15 Station Road , Crosshills,
Keighley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Web www.viamed.co.uk

COCA4P860RA

Certificate of Compatibility
Class IIb

SpO_2 probe - pulse oximetry sensor extension cable:

For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.

I the undersigned certify that the part(s) listed **P927E8**
is/are compatible with the manufacturer(s) part (s) **Spacelabs**

It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.

This declaration is supported by:

Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086

Signed: Date:25th November 2003

Position: Managing Director

VIAMED

Viamed Ltd. 15 Station Road , Crosshills,
Keigley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Web www.viamed.co.uk

Certificate of Compatibility
Class IIb

SpO_2 probe - pulse oximetry sensor:

For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.

I the undersigned certify that the part(s) listed **P858YA**
is/are compatible with the manufacturer(s) part (s) **Nellcor D-YS (Dura-Y)**

It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.

This declaration is supported by:

Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086

Signed: Date: 25 November 2003

Position: Managing Director

VIAMED
Viamed Ltd. 15 Station Road , Crosshills,
Keigley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Web www.viamed.co.uk

VIAMED

Certificate of Compatibility
Class IIb

SpO2 probe - pulse oximetry sensor:
For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.
I the undersigned certify that the part(s) listed P973E10
is/are compatible with the manufacturer(s) part (s) Datex® SAS-C3
It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.
This declaration is supported by:
Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086
Signed: Date: 23 April 1999
Position: Managing Director

Viamed Ltd. 15 Station Road , Crosshills,
Keigley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Webb www.viamed.co.uk cocP973e10

Certificate of Compatibility
Class IIb

SpO2 probe - pulse oximetry sensor:
For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.
I the undersigned certify that the part(s) listed P973E10
is/are compatible with the manufacturer(s) part (s) Datex® SAS-C3
It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.
This declaration is supported by:
Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086
Signed: Date: 23 April 1999
Position: Managing Director

Viamed Ltd. 15 Station Road , Crosshills,
Keigley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Webb www.viamed.co.uk cocP973E10
Certificate of Compatibility
Class IIb

SpO2 probe - pulse oximetry sensor:
For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.
I the undersigned certify that the part(s) listed P973E10
is/are compatible with the manufacturer(s) part (s) Datex® SAS-C3
It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.
This declaration is supported by:
Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)

Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086
Signed: Date: 23 April 1999
Position: Managing Director

Viamed Ltd. 15 Station Road , Crosshills,
Keigley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Webb www.viamed.co.uk cocP956e8

Certificate of Compatibility
Class IIb

SpO2 probe - pulse oximetry sensor:
For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.
I the undersigned certify that the part(s) listed P956E4
is/are compatible with the manufacturer(s) part (s) Nellcor® EC4
It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.
This declaration is supported by:
Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086
Signed: Date: 23 April 1999
Position: Managing Director

Viamed Ltd. 15 Station Road , Crosshills,
Keigley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Webb www.viamed.co.uk cocP956E8

Certificate of Compatibility
Class IIb

SpO2 probe - pulse oximetry sensor:

For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.

I the undersigned certify that the part(s) listed P891RA
is/are compatible with the manufacturer(s) part (s) Critikon

It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.

This declaration is supported by:

Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086

Signed: Date: 11 September 2001

Position: Managing Director

Viamed Ltd. 15 Station Road , Crosshills,
Keigley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Web www.viamed.co.uk

Certificate of Compatibility
Class IIb

SpO2 probe - pulse oximetry sensor:

For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.

I the undersigned certify that the part(s) listed P863RA
is/are compatible with the manufacturer(s) part (s) DatascopeR 0600-00-0026-01

It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.

This declaration is supported by:

Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086

Signed: Date: 23 April 1999

Position: Managing Director

Viamed Ltd. 15 Station Road , Crosshills,
Keigley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Web www.viamed.co.uk

COCA4M

Certificate of Compatibility
Class IIb

SpO_2 probe - pulse oximetry sensor extension cable:

For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.

I the undersigned certify that the part(s) listed **P973E10**
is/are compatible with the manufacturer(s) part (s) **Datex**

It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.

This declaration is supported by:

Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086

Signed: Date:25th November 2003

Position: Managing Director

VIAMED

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Keigley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Web www.viamed.co.uk

Certificate of Compatibility
Class IIb

SpO_2 probe - pulse oximetry sensor extension cable:

For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.

I the undersigned certify that the part(s) listed **P965E10**
is/are compatible with the manufacturer(s) part (s) **Critikon & Sensormedics**

It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.

This declaration is supported by:

Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086

Signed: Date:25th November 2003

Position: Managing Director

VIAMED

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Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Web www.viamed.co.uk

Certificate of Compatibility
Class IIb

SpO_2 probe - pulse oximetry sensor extension cable:

For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.

I the undersigned certify that the part(s) listed **P973E33**
is/are compatible with the manufacturer(s) part (s) **Datex**

It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.

This declaration is supported by:

Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086

Signed: Date:25th November 2003

Position: Managing Director

VIAMED

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Keigley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Web www.viamed.co.uk

Certificate of Compatibility
Class IIb

SpO_2 probe - pulse oximetry sensor extension cable:

For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.

I the undersigned certify that the part(s) listed **P925E2**
is/are compatible with the manufacturer(s) part (s) **Spacelabs**

It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.

This declaration is supported by:

Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086

Signed: Date:25th November 2003

Position: Managing Director

VIAMED

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Keigley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Web www.viamed.co.uk

[redacted]

SpO_2 probe - pulse oximetry sensor:
For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.

I the undersigned certify that the part(s) listed **P973E10**
is/are compatible with the manufacturer(s) part (s) **Datex® SAS-C3**
It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.
This declaration is supported by:
Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086
Signed: Date: 23 April 1999
Position: Managing Director

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Keigley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Webb www.viamed.co.uk COCP973e10

[redacted]

SpO_2 probe - pulse oximetry sensor:
For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.

I the undersigned certify that the part(s) listed **P973E5**
is/are compatible with the manufacturer(s) part (s) **Datex® SAS-C1**
It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.
This declaration is supported by:
Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086
Signed: Date: 23 April 1999
Position: Managing Director

Viamed Ltd. 15 Station Road , Crosshills,
Keigley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Webb www.viamed.co.uk COCP973E5

[redacted]

SpO_2 probe - pulse oximetry sensor:
For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.

I the undersigned certify that the part(s) listed **P973E10**
is/are compatible with the manufacturer(s) part (s) **Datex® SAS-C3**
It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.

This declaration is supported by:
Technical documentation required by the MDD (
EC Quality Assurance Certificate No. CE 01389(
Issued by: British Standards Institute Dat
Notified body number CE 0086
Signed: Date
Position: Managing Director

Viamed Ltd. 15 Station Road , Crosshills,
Keigley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Webb www.viamed.c

[redacted]

SpO_2 probe - pulse oximetry sensor:
For use with pulse oximeters in order to determi

I the undersigned certify that the part(s) listed **P**
is/are compatible with the manufacturer(s) part
It is also hereby declared that the medical devic
requirements listed in Annex I of the EC Counc
This declaration is supported by:
Technical documentation required by the MDD
EC Quality Assurance Certificate No. CE 0138
Issued by: British Standards Institute Da
Notified body number CE 0086
Signed: Da
Position: Managing Director

Viamed Ltd. 15 Station Road , Crosshills,
Keigley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Webb www.viamed

Certificate of Compatibility
Class IIb

SpO_2 probe - pulse oximetry sensor:

For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.

I the undersigned certify that the part(s) listed **P863RA**
is/are compatible with the manufacturer(s) part (s) **Datascope[R]** **0600-00-0026-01**

It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.

This declaration is supported by:

Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086

Signed: Date: 23 April 1999

Position: Managing Director

VIAMED

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Keigley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Web www.viamed.co.uk

Certificate of Compatibility
Class IIb

SpO_2 probe - pulse oximetry sensor:

For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.

I the undersigned certify that the part(s) listed **P875YA**
is/are compatible with the manufacturer(s) part (s) **Novametrics**

It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.

This declaration is supported by:

Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086

Signed: Date: 25 November 2003

Position: Managing Director

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Keigley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Web www.viamed.co.uk

Certificate of Compatibility
Class IIb

SpO_2 probe - pulse oximetry sensor:

For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.

I the undersigned certify that the part(s) listed **P858RA**
is/are compatible with the manufacturer(s) part (s) **Nellcor DS100A**

It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.

This declaration is supported by:

Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086

Signed: Date: 22 October 2003

Position: Managing Director

Viamed Ltd. 15 Station Road , Crosshills,
Keigley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Web www.viamed.co.uk

[redacted]

SpO_2 probe - pulse oximetry sensor:

For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.

I the undersigned certify that the part(s) listed **P877RA**
is/are compatible with the manufacturer(s) part (s) **CRITIKON®** **8997**

It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.

This declaration is supported by:

Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086

Signed: Date: 23 April 1999

Position: Managing Director

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Keighley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Web www.viamed.co.uk

[redacted]

SpO_2 probe - pulse oximetry sensor:

For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.

I the undersigned certify that the part(s) listed **P876YA**
is/are compatible with the manufacturer(s) part (s) **Novametrics®** **Flexible probe**

It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.

This declaration is supported by:

Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086

Signed: Date: 23 April 1999

Position: Managing Director

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Keighley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Web www.viamed.co.uk

[redacted]

SpO_2 probe - pulse oximetry sensor:

For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.

I the undersigned certify that the part(s) listed **P876RA**
is/are compatible with the manufacturer(s) part (s) **Novametrics®** **8774**

It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.

This declaration is supported by:

Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086

Signed: Date: 23 April 1999

Position: Managing Director

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Keighley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Web www.viamed.co.uk

[redacted]

SpO_2 probe - pulse oximetry sensor:

For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.

I the undersigned certify that the part(s) listed **P873RA**
is/are compatible with the manufacturer(s) part (s) **DATEX®** **SAS-F4**

It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.

This declaration is supported by:

Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086

Signed: Date: 23 April 1999

Position: Managing Director

Viamed Ltd. 15 Station Road , Crosshills,
Keighley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Web www.viamed.co.uk

[redacted]

SpO_2 probe - pulse oximetry sensor:

For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.

I the undersigned certify that the part(s) listed **P873YA**
is/are compatible with the manufacturer(s) part (s) **DATEX®** **Flexible probe**

It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.

This declaration is supported by:

Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086

Signed: Date: 23 April 1999

Position: Managing Director

Viamed Ltd. 15 Station Road , Crosshills,
Keighley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Web www.viamed.co.uk

SpO_2 probe - pulse oximetry sensor:

For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.

I the undersigned certify that the part(s) listed **P866RA**
is/are compatible with the manufacturer(s) part (s) **Simed ®/ Finger probes**

It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.

This declaration is supported by:

Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086

Signed: Date: 23 April 1999

Position: Managing Director

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Keighley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Web www.viamed.co.uk

SpO_2 probe - pulse oximetry sensor:

For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.

I the undersigned certify that the part(s) listed **P865RA**
is/are compatible with the manufacturer(s) part (s) **Sensormedics ®/Critikon ®** **Finger probes**

It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.

This declaration is supported by:

Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086

Signed: Date: 23 April 1999

Position: Managing Director

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Keighley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Web www.viamed.co.uk

[redacted]

SpO_2 probe - pulse oximetry sensor:

For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.

I the undersigned certify that the part(s) listed **P856YA**
is/are compatible with the manufacturer(s) part (s) **NELLCOR®** **Dura Y ®**

It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.

This declaration is supported by:

Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086

Signed: Date: 23 April 1999

Position: Managing Director

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Keighley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Web www.viamed.co.uk

[redacted]

SpO_2 probe - pulse oximetry sensor:

For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.

I the undersigned certify that the part(s) listed **P872RA**
is/are compatible with the manufacturer(s) part (s) **DATEX®** **SAS-F4**

It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.

This declaration is supported by:

Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086

Signed: Date: 23 April 1999

Position: Managing Director

Viamed Ltd. 15 Station Road , Crosshills,
Keighley, West Yorkshire, BD20 7DT. UK.
Tel +44 1535 634542. Fax +44 1535 635582
email info@viamed.co.uk Web www.viamed.co.uk

SpO_2 probe - pulse oximetry sensor:

For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.

I the undersigned certify that the part(s) listed **P864RA**
is/are compatible with the manufacturer(s) part (s) **DATASCOPE®** **0600-00-0026-02**

It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.

This declaration is supported by:

Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
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Position: Managing Director

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[redacted]

SpO_2 probe - pulse oximetry sensor:

For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.

I the undersigned certify that the part(s) listed **P856RA**
is/are compatible with the manufacturer(s) part (s) **NELLCOR®** **DS-100A**

It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.

This declaration is supported by:

Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086

Signed: Date: 23 April 1999

Position: Managing Director

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[redacted]

SpO_2 probe - pulse oximetry sensor:

For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.

I the undersigned certify that the part(s) listed **PXN**
is/are compatible with the manufacturer(s) part (s) **OEM ®** **OEMNO**

It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.

This declaration is supported by:

Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086

Signed: Date: 23 April 1999

Position: Managing Director

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COCA4M

Certificate of Compatibility
Class IIb

SpO_2 probe - pulse oximetry sensor extension cable:

For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively.

I the undersigned certify that the part(s) listed **P971E8**
is/are compatible with the manufacturer(s) part (s) **Nonin**

It is also hereby declared that the medical devices specified above conforms with the essential requirements listed in Annex I of the EC Council Directive 93/42/EEC of 14 June 1993.

This declaration is supported by:

Technical documentation required by the MDD (Annex VII) retained by: Viamed Limited
EC Quality Assurance Certificate No. CE 01389(Annex V)
Issued by: British Standards Institute Date: 8 February 1999
Notified body number CE 0086

Signed: Date:25th November 2003

Position: Managing Director

VIAMED

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