



CERTIFICATE OF ASSESSMENT - EC

DET NORSKE VERITAS

Application of Council Directive 93/42/EEC of 14 June 1993, adopted as "Forskrift for Medisinsk Utstyr" by the Norwegian Ministry of Health and Social Affairs.

Certificate N°.: 2001-OSL-MDD-0174

This is to certify that the Quality System for the product group:

Active Therapeutic Device

- defined by manufacturer as Class IIa devices -

Manufactured by
MULTI-WELL

Z.I. du Grand Pont BP 46
F-13640 LA ROQUE D'ANTHERON, FRANCE

complies with the applicable requirements of the Directive.

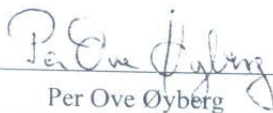
The quality system for these products has been assessed according to the procedure of conformity assessment described in **Article 11.2.b) and Annex V**. Identification of the products covered by this certificate is given in the Appendix.

Limitations:

The manufacturer must inform Det Norske Veritas Region Norge AS of any plan for substantial changes to the quality system in order to examine whether this Certificate remains valid. Annual Periodical Audits will be held to verify the validity of this Certificate.

Høvik, 26 June 2001

for Det Norske Veritas Region Norge



Per Ove Øyberg
Head of section, Testing, Product and Personnel
Certification

CE
0434

Valid until: 26 June 2006



Eugenie Winger Husebye
Service responsible Medical Devices

This Certificate is valid until the date specified. Any significant changes in the design or construction of the products, the quality system or amendments to the Directive may render this Certificate invalid at an earlier date. The product liability rests with the manufacturer or his representative in accordance with Council Directive 85/374/EEC



CERTIFICATE OF ASSESSMENT - EC

APPENDIX



Appendix to Certificate No.: 2001-OSL-MDD-0174
Manufacturer: MULTI-WELL
Product group: Active Therapeutic Device

The Certificate referred to above covers the following devices:

- Bioderm 40

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26 June 2001

Eugenie Winger Husebye
for Eugenie Winger Husebye

CE0434



CERTIFICATE OF ASSESSMENT - EC

DET NORSKE VERITAS

Application of Council Directive 93/42/EEC of 14 June 1993, adopted as "Forskrift for Medisinsk Utstyr" by the Norwegian Ministry of Health and Social Affairs.

Certificate N°.: 2005-OSL-MDD-0145

This is to certify that the Quality System for the product group:

Ultra-sonic waves emission devices for therapeutic use

- defined by manufacturer as Class IIa devices -

Manufactured by

MULTI-WELL

Z.I. du Grand Pont BP 46 F-13640 LA ROQUE D'ANTHERON, FRANCE

complies with the applicable requirements of the Directive.

The quality system for these products has been assessed according to the procedure of conformity assessment described in **Article 11.3.a) and Annex II (excl. section 4)**. Identification of the products covered by this certificate is given in the Appendix.

Limitations:

The manufacturer must inform Det Norske Veritas Certification AS of any plan for significant changes to the quality system. Annual Periodical Audits will be held to verify the validity of this Certificate.

Hovik, 21 January 2005

for Det Norske Veritas Certification AS

Line Gangeskar

Line Gangeskar

Head of section, Testing, Product and Personnel
Certification

CE
0434

Valid until: 21 January 2010

Eugenie Winger Husebye

Eugenie Winger Husebye
Service responsible Medical Devices

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CERTIFICATE OF ASSESSMENT - EC APPENDIX


Appendix to Certificate No.: 2005-OSL-MDD-0145
Manufacturer: MULTI-WELL
Product group: Ultra-sonic waves emission devices for therapeutic use

The Certificate referred to above covers the following devices:

- **Cell Sonic NG**
 - **Models: Multiwell SUP, Multiwell BAS, Algobody, Ultrasons**

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20 April 2005


Eugenie Winger Husebye





CERTIFICATE OF ASSESSMENT - EC

DET NORSKE VERITAS

Application of Council Directive 93/42/EEC of 14 June 1993, adopted as "Forskrift for Medisinsk Utstyr" by the Norwegian Ministry of Health and Social Affairs.

Certificate N°.: 2005-OSL-MDD-0144

This is to certify that the Quality System for the product group:

Pressure -Therapy devices

- defined by manufacturer as Class IIa devices -

Manufactured by

MULTI-WELL

Z.I. du Grand Pont BP 46 F-13640 LA ROQUE D'ANTHERON, FRANCE

complies with the applicable requirements of the Directive.

The quality system for these products has been assessed according to the procedure of conformity assessment described in **Article 11.3.a) and Annex II (excl. section 4)**. Identification of the products covered by this certificate is given in the Appendix.

Limitations:

The manufacturer must inform Det Norske Veritas Certification AS of any plan for significant changes to the quality system. Annual Periodical Audits will be held to verify the validity of this Certificate.

Høvik, 21 January 2005

for Det Norske Veritas Certification AS

Line Gangeskar

Line Gangeskar
*Head of section, Testing, Product and Personnel
Certification*

CE
0434

Valid until: 21 January 2010

Eugenie Winger Husebye

Eugenie Winger Husebye
Service responsible Medical Devices

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CERTIFICATE OF ASSESSMENT - EC APPENDIX

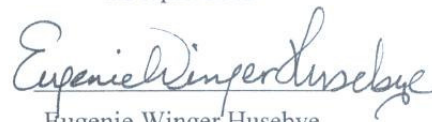
Appendix to Certificate No.: 2005-OSL-MDD-0144
Manufacturer: MULTI-WELL
Product group: Pressure -Therapy devices

The Certificate referred to above covers the following devices:

- **DLS 10 NG**
 - **Models: Algopress, Presso, DLS 10 S103 Module SUP, DLS 10 S103 Module BAS, DLS 10 S102 Module SUP, DLS 10 S102 Module BAS**

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20 April 2005


Eugenie Winger Husebye





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DET NORSKE VERITAS MANAGEMENT SYSTEM CERTIFICATE

Certificate No. 99-OSL-AQ-6839

This is to certify that
THE QUALITY SYSTEM
of

MultiWell

at
Z.I. du Grand Pont
F-13640 LA ROQUE d'ANTHERON
FRANCE

has been found to conform to the Quality System Standard
NS-EN ISO 9001 : 2000 / ISO 13485 : 1996

This Certificate is valid for the following product or service ranges:

**The development, assembly and sales of measuring, investigational and treatment equipment used for medical, para-medical and sportive applications and the sales of applied software.
Development and sales of diet, nutritional and cosmetic products.**

Original certificate valid from:
1999-09-14

This Certificate with Appendix is valid until:
2006-09-09

Sterling Kwarts
Lead Auditor



Place and date:
Høvik, 2003-11-17

For the accredited unit:
Det Norske Veritas Certification AS

Eugenie Winger Husebye
Management Representative

Lack of fulfilment of conditions as set out in the Appendix may render this Certificate invalid.



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Application of Council Directive 93/42/EEC of 14 June 1993, adopted as "Forskrift for Medisinsk Utstyr" by the Norwegian Ministry of Health and Social Affairs.

Certificate N°.: 2005-OSL-MDD-0146

This is to certify that the Quality System for the product group:

Physiologic parameters measures by bio impedance devices

- defined by manufacturer as Class IIa devices -

Manufactured by

MULTI-WELL

Z.I. du Grand Pont BP 46 F-13640 LA ROQUE D'ANTHERON, FRANCE

complies with the applicable requirements of the Directive.

The quality system for these products has been assessed according to the procedure of conformity assessment described in **Article 11.3.a) and Annex II (excl. section 4)**. Identification of the products covered by this certificate is given in the Appendix.

Limitations:

The manufacturer must inform Det Norske Veritas Certification AS of any plan for significant changes to the quality system. Annual Periodical Audits will be held to verify the validity of this Certificate.

Høvik, 21 January 2005

for Det Norske Veritas Certification AS

Line Gangeskar

Line Gangeskar

*Head of section, Testing, Product and Personnel
Certification*

CE
0434

Valid until: 21 January 2010

Eugenie Winger Husebye

Eugenie Winger Husebye
Service responsible Medical Devices

This Certificate is valid until the date specified. Any significant changes in the design or construction of the products, the quality system or amendments to the Directive may render this Certificate invalid at an earlier date. The product liability rests with the manufacturer or his representative in accordance with Council Directive 85/374/EEC



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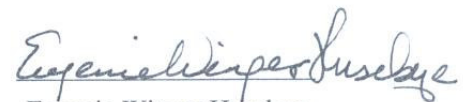
Appendix to Certificate No.: 2005-OSL-MDD-0146
Manufacturer: MULTI-WELL
Product group: Physiologic parameters measures by bio impedance devices

The Certificate referred to above covers the following devices:

➤ Ohmniscan NG

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20 April 2005


Eugenie Winger Husebye

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