



Declaration of Conformity

Certificate Number:

LTE Scientific Limited, hereby declare that the pressure equipment specified below has been designed, manufactured and tested in accordance with the requirements of the Pressure Equipment Regulations 1999. (European Directive 97/23/EC).
The Electromagnetic Compatibility Directive (2004/108/EC)
The Low Voltage Directive (2006/95/EC)

Description of Equipment

Product Type: **Non Porous Load Autoclave** Model: TOUCHCLAVE – LAB
Serial Number: PV0 Date of Manufacture:

Component	Classification	Procedure Followed	EC Design Examination Number
Vessel Chamber	Category	Module	CE-41-PED-B-DFP-001-03-GBR
Steam Generator	Category	Module	
Piping and Pressure Accessories	SEP	N/A	
Safety Relief Valves	Category IV	Module B+D	

Name of Notified Body for Product Verification: **HSB INSPECTION QUALITY LTD**

Notified Body Reference: **0878**

Address of Notified Body: **Cairo House Greenacres Road Waterhead Oldham England OL4 3JA**

Applied Harmonised Standards: **None Applied.**
Other Standards Applied: **PD5500**
EN61010-1 2001
EN55011
EN61000-4

Note: This product has been manufactured and tested to satisfy the requirements of your purchase order and our acknowledgement of your order. Under no circumstances must it be used for any other purpose. The product must be installed and maintained by a competent person in accordance with the user manual provided with the equipment.

Authorised Signatory for LTE SCIENTIFIC LTD.

Name: Ian J Dust Position: Quality Supervisor

CE
0086

Signature: _____

Date:



Registered No. 2648370 (England)

LTE SCIENTIFIC LTD
Greenbridge Lane Greenfield Oldham United Kingdom OL3 7EN

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E: info@lte-scientific.com W: www.lte-scientific.com



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Description of Equipment

Product Type: **Laboratory Autoclave** Model: TCR
 Serial Number: PV0 Date of Manufacture:

Component	Classification	Procedure Followed	EC Design Examination Number
Vessel Chamber	Category	Module	
Steam Generator	Category	Module	
Piping and Pressure Accessories	SEP	N/A	
Safety Relief Valves	Category IV	Module B+D	

Name of Notified Body for Product Verification: **HSB INSPECTION QUALITY LTD**
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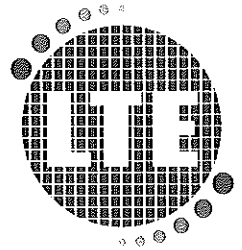


Name: Ian J Dust Position: Quality Supervisor

Signature: _____

Date:





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Certificate Number:

LTE Scientific Limited, hereby declare that the pressure equipment specified below has been designed, manufactured and tested in accordance with the requirements of the Pressure Equipment Directive (97/23/EC)

Medical Device Directive (93/42/EC)

Description of Equipment

Product Type: **Porous Load Autoclave** Model: TOUCHCLAVE - PL
 Serial Number: PVO Date of Manufacture:

Component	Classification	Procedure Followed	EC Design Examination Number
Vessel Chamber	Category	Module	CE-41-PED-B-DFP-001-03-GBR
Steam Generator	Category	Module	
Piping and Pressure Accessories	SEP	N/A	
Safety Relief Valves	Category IV	Module B+D	

Name of Notified Body for Product Verification: **HSB INSPECTION QUALITY LTD**

Notified Body Reference: **0878**

Address of Notified Body: **Cairo House Greenacres Road Waterhead Oldham England OL4 3JA**

Applied Harmonised Standards: **None Applied.**
 Other Standards Applied: **PD5500**
EN61010-1 2001
EN55011
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Name: Ian J Dust Position: Quality Supervisor

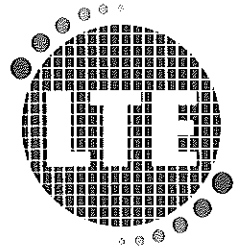
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Medical Device Directive (93/42/EC)

Description of Equipment

Product Type: **Instrument Autoclave**

Model: TCR

Serial Number: PVO

Date of Manufacture:

Component	Classification	Procedure Followed	EC Design Examination Number
Vessel Chamber	Category	Module	
Steam Generator	Category	Module	
Piping and Pressure Accessories	SEP	N/A	
Safety Relief Valves	Category IV	Module B+D	

Name of Notified Body for Product Verification: **HSB INSPECTION QUALITY LTD**

Notified Body Reference: **0878**

Address of Notified Body: **Cairo House Greenacres Road Waterhead Oldham England OL4 3JA**

Applied Harmonised Standards:

None Applied.

Other Standards Applied:

**PD5500
EN61010-1 2001
EN55011
EN61000-4**

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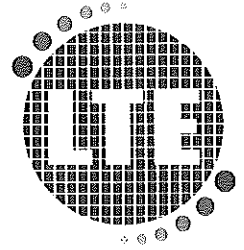
Name: Ian J Dust

Position: Quality Supervisor

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Date:





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LTE Scientific Limited, hereby declare that the pressure equipment specified below has been designed, manufactured and tested in accordance with the requirements of the Pressure Equipment Directive (97/23/EC)
Medical Device Directive (93/42/EC)

Description of Equipment

Product Type: **Touchclave Systems Non Porous Load Autoclaves Single and Double Door Chamber Sizes 65 Litre to 4000 Litre**

Model: Serial Number: PV0 Date of Manufacture:

Component	Classification	Procedure Followed	EC Design Examination Number
Vessel Chamber	Category	Module	F Certificate available
Piping and Pressure Accessories	SEP	N/A	
Safety Relief Valves	Category IV	Module B+D	CE-41-PED-B-DFP-001-03-GBR
Interlock and Assemblies		Module B+D	SS 42070682-0002 SS 42070682-0003 CEN-019596

Name of Notified Body for Product Verification: **Zurich Risk Services**

Notified Body Reference: **0037**

Address of Notified Body: **Park House Bristol Road South Rubery Birmingham England B45 9AH**

Applied Harmonised Standards:	None Applied.	
Other Standards Applied:	PD5500	
	WUDT/UC/2003	EN61010-1 2001
	EN554 1994	EN55011
	EN556 1995	EN285 1997
	EN61000-4	

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CE
0086 0037

Name: John Cruse Position: Technical Manager

Signature: _____

Date:



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Description of Equipment

Product Type: **Touchclave Systems Non Porous Load Autoclaves Single and Double Door Chamber Sizes 65 Litre to 4000 Litre**

Model: Serial Number: PV0 Date of Manufacture:

Component	Classification	Procedure Followed	EC Design Examination Number
Vessel Chamber	Category	Module	F Certificate Available
Piping and Pressure Accessories	SEP	N/A	
Safety Relief Valves	Category IV	Module B+D	CE-41-PED-B-DFP-001-03-GBR
Interlock and Assemblies		Module B+D	SS 42070682-0002 SS 42070682-0003 CEN-019596

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Applied Harmonised Standards:	None Applied.	
Other Standards Applied:	PD5500	
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Name: John Cruse Position: Technical Manager

Signature: _____

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