STATUTORY RULES OF NORTHERN IRELAND

2003 No. 386

HEALTH AND SAFETY

Transportable Pressure Vessels Regulations (Northern Ireland) 2003

Made - - - - 22nd August 2003

Coming into operation –

regulations 1, 2, 10 and 12(1) to (5)

1st October 2003

remaining regulations

1st December 2003

TRANSPORTABLE PRESSURE VESSELS REGULATIONS (NORTHERN IRELAND) 2003

PART 1

PRELIMINARY

- 1. Citation and commencement
- 2. Interpretation
- 3. Application

PART 2

GENERAL REQUIREMENTS

- 4. Requirements relating to the placing on the market and use at work of transportable pressure vessels
- 5. Transportable pressure vessels placed on the market or used at work exclusively in Northern Ireland
- 6. Reassessment of conformity
- 7. Periodic inspection and repeated use
- 8. Notified bodies
- 9. Approved bodies
- 10. Appointment of notified bodies and approved bodies by the Executive
- 11. Conformity marking

PART 3

MISCELLANEOUS

1	
	Fees

- 13. Defence
- 14. Amendments and saving Signature

SCHEDULE DISAPPLICATIONS TO THESE REGULATIONS

1

- 1. These Regulations shall not apply to –
- 2. These Regulations shall not apply to any transportable pressure vessel...
- 3. These Regulations shall not apply to transportable pressure vessels

-...

SCHEDULE STANDARDS

2

SCHEDULE CONFORMITY ASSESSMENT PROCEDURES

3

(This Schedule substantially reproduces the provisions of Part 1 of Annex IV to the Transportable Pressure Equipment Directive.)

- 1. Module A—internal production control
- 2. The manufacturer must draw up the technical documentation described in...
- 3. The technical documentation must enable an assessment to be made...
- 4. The manufacturer, or his authorised representative established within the Community,...
- 5. The manufacturer must take all measures necessary to ensure that... Module A1—internal manufacturing checks with monitoring of the final assessment
- 1. Module B—EC type-examination
- 2. The application for EC-type-examination must be lodged by the manufacturer...
- 3. The technical documentation must enable an assessment to be made...
- 4. The notified body must:
- 4.1 examine the technical documentation, verify that the type has been...
- 4.2 perform or have performed the appropriate examinations and necessary tests...
- 4.3 perform or have performed the appropriate examinations and necessary tests...
- 4.4 agree with the applicant the location where the examinations and...
- 5. Where the type satisfies the relevant provisions of these Regulations,...
- 6. The applicant must inform the notified body that holds the...
- 7. Each notified body must communicate to the Member States the...
- 8. The other notified bodies may receive copies of the EC...

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- 9. The manufacturer, or his authorised representative established within the Community,...
- 1. Module B1—EC design examination
- 2. The manufacturer, or his authorised representative established within the Community,...
- 3. The technical documentation must enable an assessment to be made...
- 4. The notified body must:
- 4.1 examine the technical documentation and identify the components which have...
- 4.2 perform the necessary examinations to establish whether the solutions adopted...
- 4.3 perform the necessary examinations to establish whether the relevant provisions...
- 5. Where the design meets the provisions of these Regulations which...
- 6. The applicant must inform the notified body that holds the...
- 7. Each notified body must communicate to the member States the...
- 8. The other notified bodies may on request obtain the relevant...
- 9. The manufacturer, or his authorised representative established within the Community,...
- 1. Module C1—conformity to type
- 2. The manufacturer must take all measures necessary to ensure that...
- 3. The manufacturer, or his authorised representative established within the Community,...
- 4. Final assessment must be subject to monitoring in the form...
- 1. Module D—production quality assurance
- 2. The manufacturer must operate an approved quality system for production,...
- 3. Quality system
- 3.1 The manufacturer must lodge an application for assessment of his...
- 3.2 The quality system must ensure compliance of the transportable pressure...
- 3.3 The notified body must assess the quality system to determine...
- 3.4 The manufacturer must undertake to fulfil the obligations arising out...
- 4. Surveillance under the responsibility of the notified body
- 4.1 The purpose of surveillance is to ensure that the manufacturer...
- 4.2 The manufacturer must allow the notified body access for inspection...
- 4.3 The notified body must carry out periodic audits to make...
- 4.4 In addition, the notified body may pay unexpected visits to...
- 5. The manufacturer must, for a period of 10 years after...
- 6. Each notified body must communicate to the member States the...
- 1. Module D1—production quality assurance
- 2. The manufacturer must draw up the technical documentation described below....
- 3. The manufacturer must operate an approved quality system for production,...
- 4. Quality system
- 4.1 The manufacturer must lodge an application for assessment of his...
- 4.2 The quality system must ensure compliance of the transportable pressure...
- 4.3 The notified body must assess the quality system to determine...

- 4.4 The manufacturer must undertake to fulfil the obligations arising out...
- 5. Surveillance under the responsibility of the notified body
- 5.1 The purpose of surveillance is to make sure that the...
- 5.2 The manufacturer must allow the notified body access for inspection...
- 5.3 The notified body must carry out periodic audits to make...
- 5.4 In addition, the notified body may pay unexpected visits to...
- 6. The manufacturer must, for a period of 10 years after...
- 7. Each notified body must communicate to the member States the...
- 1. Module E—product quality assurance
- 2. The manufacturer must operate an approved quality system for production,...
- 3. Quality system
- 3.1 The manufacturer must lodge an application for assessment of his...
- 3.2 Under the quality system, each transportable pressure vessel must be...
- 3.3 The notified body must assess the quality system to determine...
- 3.4 The manufacturer must undertake to fulfil the obligations arising out...
- 4. Surveillance under the responsibility of the notified body
- 4.1 The purpose of surveillance is to ensure that the manufacturer...
- 4.2 The manufacturer must allow the notified body access for inspection...
- 4.3 The notified body must carry out periodic audits to make...
- 4.4 In addition, the notified body may pay unexpected visits to...
- 5. The manufacturer must, for a period of 10 years after...
- 6. Each notified body must communicate to the member States the...
- 1. Module E1—production quality assurance
- 2. The manufacturer must draw up the technical documentation described below....
- 3. The manufacturer must operate an approved quality system for the...
- 4. Quality system
- 4.1 The manufacturer must lodge an application for assessment of his...
- 4.2 Under the quality system, each transportable pressure vessel must be...
- 4.3 The notified body must assess the quality system to determine...
- 4.4 The manufacturer must undertake to discharge the obligations arising from...
- 5. Surveillance under the responsibility of the notified body
- 5.1 The purpose of surveillance is to ensure that the manufacturer...
- 5.2 The manufacturer must allow the notified body access for inspection...
- 5.3 The notified body must carry out periodic audits to make...
- 5.4 In addition, the notified body may pay unexpected visits to...
- 6. The manufacturer must, for a period of 10 years after...
- 7. Each notified body must communicate to the member States the...
- 1. Module F—product verification
- 2. The manufacturer must take all measures necessary to ensure that...
- 3. The notified body must perform the appropriate examinations and tests
- 4. Verification by examination and testing of each transportable pressure vessel

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- 4.1 Each transportable pressure vessel must be individually examined and must...
- 4.2 The notified body must affix its identification number or have...
- 4.3 The manufacturer, or his authorised representative established within the Community,...
 - 1. Module G—EC unit verification
 - 2. The manufacturer must apply to a notified body of his...
- 3. The technical documentation must enable the conformity of the transportable...
- 4. The notified body must examine the design and construction of...
- 4.1 The notified body must affix its identification number or have...
- 4.2 The manufacturer, or his authorised representative established within the Community,...
- 1. Module H—full quality assurance
- 2. The manufacturer must implement an approved quality system for design,...
- 3. Quality system
- 3.1 The manufacturer must lodge an application for assessment of his...
- 3.2 The quality system must ensure compliance of the transportable pressure...
- 3.3 The notified body must assess the quality system to determine...
- 3.4 The manufacturer must undertake to fulfil the obligations arising out...
- 4. Surveillance under the responsibility of the notified body
- 4.1 The purpose of this surveillance is to ensure that the...
- 4.2 The manufacturer must allow the notified body access for inspection...
- 4.3 The notified body must carry out periodic audits to make...
- 4.4 In addition, the notified body may pay unexpected visits to...
- 5. The manufacturer must, for a period of 10 years after...
- 6. Each notified body must communicate to the member States the...
- 1. Module H1—full quality assurance with design examination and special surveillance of the final test
- 2. Final assessment is subject to increased surveillance in the form...

SCHEDULE MODULES TO BE FOLLOWED FOR CONFORMITY

4 ASSESSMENT

(This Schedule substantially reproduces the provisions of Annex V to the Transportable Pressure Equipment Directive.)

SCHEDULE CONFORMITY REASSESSMENT PROCEDURE

5

(This Schedule substantially reproduces the provisions of Part II of Annex IV to the Transportable Pressure Equipment Directive.)

- 1. This procedure describes the method for ensuring that transportable pressure...
- 2. The owner must make available to a notified body information...
- 3. The notified body must check whether transportable pressure vessels which...
- 4. If the results of the above checks are satisfactory, the...
- 5. For vessels manufactured in series, including their valves and other...

SCHEDULE PERIODIC INSPECTION PROCEDURES

6

(This Schedule substantially reproduces the provisions of Part III of Annex IV to the Transportable Pressure Equipment Directive.)

- 1. Module 1—periodic inspection of products
- 2. To meet the requirements referred to in paragraph 1, the...
- 3. The notified or approved body must perform the appropriate examinations...
- 3.1 All transportable pressure vessels must be examined individually and appropriate...
- 3.2 The notified or approved body must affix, or have affixed,...
- 3.3 The owner or his authorised representative established in the Community...
- 1. Module 2—periodic inspection through quality assurance
- 2. The owner or his authorised representative established in the Community...
- 3. Quality system
- 3.1 The owner or his authorised representative established in the Community...
- 3.2 Under the quality system, each transportable pressure vessel must be...
- 3.3 The notified body must assess the quality system to determine...
- 3.4 The owner or his authorised representative established in the Community...
- 4. Surveillance under the responsibility of the notified body
- 4.1 The purpose of surveillance is to make sure that the...
- 4.2 The owner or his authorised representative established in the Community...
- 4.3 The notified body must carry out periodic audits to make...
- 4.4 In addition, the notified body may pay unannounced visits to...
- 5. The owner or his authorised representative established in the Community...

SCHEDULE CONFORMITY MARKING

7

(This Schedule substantially reproduces the provisions of Annex VII to the Transportable Pressure Equipment Directive.)

SCHEDULE AMENDMENTS TO THE CDGCPL REGULATIONS

8

- 1. The CDGCPL Regulations shall be amended in accordance with paragraphs...
- 2. In regulation 2(1) (a) for the definition of "competent...
- 3. For paragraphs (4) and (5) of regulation 3, there shall...
- 4. Regulations 12 to 17 shall be deleted.
- 5. In regulation 19, for paragraph (2) there shall be substituted...
- 6. In Schedule 4 (a) in paragraph 1, for the...
- 7. For Schedule 8 there shall be substituted the following Schedule...
- 8. In Schedule 9 (a) in paragraph 2, for the... Explanatory Note