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

PART 2

GENERAL REQUIREMENTS

PART 3

MISCELLANEOUS

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

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***

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 4 MODULES TO BE FOLLOWED FOR CONFORMITY ASSESSMENT

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

SCHEDULE 5 CONFORMITY REASSESSMENT PROCEDURE

(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 6 PERIODIC INSPECTION PROCEDURES

(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 7 CONFORMITY MARKING

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

SCHEDULE 8 AMENDMENTS TO THE CDGCPL REGULATIONS

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