STATUTORY INSTRUMENTS

2004 No. 568

HEALTH AND SAFETY

The Carriage of Dangerous Goods and Use of Transportable Pressure Equipment Regulations 2004

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THE CARRIAGE OF DANGEROUS GOODS AND USE OF TRANSPORTABLE PRESSURE EQUIPMENT REGULATIONS 2004

PART 1

INTRODUCTORY PROVISIONS

PART 2

REQUIREMENTS OF ADR AND RID

1. The references referred to in regulation 27(b) are–
2. Any approval granted by the GB competent authority by reference...

PART 2 — MULTILATERAL AND UNILATERAL APPROVALS BY THE GB COMPETENT AUTHORITY

3. The references referred to in regulation 28(1) are for–
4. The references referred to in regulation 28(3) in relation to...
5. (1) Where the GB competent authority grants approvals under regulation...

PART 3 — APPOINTMENT OF PERSONS BY THE GB COMPETENT AUTHORITY

6. The references referred to in regulation 29(1)(b) are–

PART 4 — RECOGNITION OF APPROVALS, TESTS, METHODS, STANDARDS, PROCEDURES ETC. BY THE GB COMPETENT AUTHORITY

7. The references referred to in regulation 30(1)(b) are—
8. (1) Where the GB competent authority recognises a quality assurance...

PART 5 — IMPOSING OF REQUIREMENTS BY THE GB COMPETENT AUTHORITY

9. The references referred to in regulation 31(1)(b) are—
10. (1) Where the GB competent authority imposes requirements by reference...

SCHEDULE 4 — CONFORMITY ASSESSMENT PROCEDURES

(This Schedule substantially reproduces the provisions of Part 1 of Annex IV of 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 of EC type-examination must be lodged by the...
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 Part 4...
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...
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 components which have been...
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 relevant provisions of Part 4...
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...
5. During such visits, the notified body must: ensure that the...
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...
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 the surveillance is to make sure that...
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 item of transportable pressure equipment...
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 make sure 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 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 item of transportable pressure equipment...
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 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 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 item of transportable...
4.1 Each item of transportable pressure equipment must be individually examined...
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...

SCHEDULE 5 — MODULES TO BE FOLLOWED FOR CONFORMITY ASSESSMENT

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

1. Transportable pressure equipment must be subject, at the choice of...
2. As part of the quality assurance procedures, the notified body...

SCHEDULE 6 — 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 equipment which...
4. If the results of the above checks are satisfactory, the...
5. For equipment manufactured in series, including their valves and other...

SCHEDULE 7 — 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 body or approved body must perform the appropriate...
3.1 All transportable pressure equipment must be examined individually and appropriate...
3.2 The notified body or approved body must affix, or have...
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 within the Community...
3. Quality system
3.1 The owner or his authorised representative established in the Community...
3.2 Under the quality system, each item of transportable pressure equipment...
3.3 The notified body must assess the quality system to determine...

6. In regulation 10 (labelling generally), in paragraph (1), for “6,...
7. In regulation 12A, in paragraph (a) for “3(2)(b), 6 or...
8. Omit Schedules 1 and 2.
9. In Schedule 3 (labels)— (1) omit paragraphs 1 to 11;...

SCHEDULE 13 — CONSEQUENTIAL AMENDMENTS

1. Amendment to the Petroleum (Consolidation) Act 1928
2. Amendment to the Compressed Acetylene (Importation) Regulations 1978
3. Amendments to the Dangerous Substances in Harbour Areas Regulations 1987
4. Amendment to the Dangerous Substances (Notification and Marking of Sites) Regulations 1990
5. Amendment to the Control of Explosives Regulations 1991
6. Amendment to the Coal and Other Safety-Lamp Mines (Explosives) Regulations 1993
7. Amendment to the Notification of New Substances Regulations 1993
8. Amendments to the Reporting of Injuries, Diseases and Dangerous Occurrences Regulations 1995
9. Amendment to the Health and Safety (Safety Signs and Signals) Regulations 1996
10. Amendments to the Pressure Systems Safety Regulations 2000
11. Amendments to the Radiation (Emergency Preparedness and Public Information) Regulations 2001
12. Amendments to the Control of Asbestos at Work Regulations 2002
13. Amendment to the Control of Lead at Work Regulations 2002
14. Amendment to the Control of Substances Hazardous to Health Regulations 2002
15. Amendment to the Dangerous Substances and Explosive Atmospheres Regulations 2002

SCHEDULE 14 — REVOCATIONS

Explanatory Note