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

- 1. Citation and commencement
- 2. Interpretation
- 3. Application
- 4. Application to international carriage
- 5. Application to tanks, pressure receptacles, battery-vehicles, battery-wagons, MEGCs, UN-certified MEGCs and transportable pressure equipment
- 6. Application to armed forces
- 7. Exemptions
- 8. Competent authority

PART 2

REQUIREMENTS OF ADR AND RID

- 9. Training
- 10. Safety obligations
- 11. Class 7 goods for carriage by rail
- 12. Safety advisers
- 13. Reports on accidents or incidents
- 14. Emergency plans for marshalling yards
- 15. Classification of goods
- 16. Prohibition from carriage
- 17. Dangerous goods list and special provisions

Status: This is the original version (as it was originally made).

- 18. Use of packagings
- 19. Use of tanks, battery-vehicles, battery-wagons, MEGCs and UN-certified MEGCs
- 20. Consignment
- 21. Construction and testing of packagings and packages
- 22. Construction and testing of tanks, battery-vehicles, battery-wagons, MEGCs and UN-certified MEGCs
- 23. Carriage, loading, unloading and handling
- 24. Vehicle crews, equipment, operation and documentation
- 25. Construction and approval of vehicles

PART 3

COMPETENT AUTHORITY FUNCTIONS

- 26. Interpretation of Part 3 and Schedule 3
- 27. Grant of approvals by the GB competent authority
- 28. Grant of unilateral and multilateral approvals by the GB competent authority in relation to class 7 goods for carriage by rail
- 29. Appointment of persons by the GB competent authority
- 30. Recognition of approvals, tests, methods, standards and procedures etc. by the GB competent authority
- 31. Imposing of requirements by the GB competent authority
- 32. Issuing of safety adviser vocational training certificates by the GB competent authority
- 33. Issuing of driver training certificates by the GB competent authority
- 34. Notification under sub-section 1.8.2.2 of ADR or of RID
- 35. Miscellaneous functions of the GB competent authority
- 36. Exemption certificates, temporary and ad hoc exemptions

PART 4

TRANSPORTABLE PRESSURE EQUIPMENT

- 37. Interpretation of Part 4
- 38. Placing on the market and use at work of transportable pressure equipment
- 39. Transportable pressure equipment placed on the market and used at work exclusively in Great Britain
- 40. Reassessment of conformity
- 41. Periodic inspection and repeated use
- 42. Notified bodies
- 43. Approved bodies
- 44. Appointment of notified bodies and approved bodies by the competent authority
- 45. Conformity marking

PART 5

ADDITIONAL REQUIREMENTS TO ADR AND RID

- 46. Attendant for carriage of class 1 goods by road
- 47. Duration of carriage and delivery of class 1 goods by road
- 48. Miscellaneous security requirements for carriage of class 1 goods by road
- 49. Miscellaneous security requirements for carriage of class 1 goods by rail
- 50. Security requirement for carriage of class 1 goods by road or rail and class 7 goods by rail

Status: This is the original version (as it was originally made).

- 51. Carriage of class 1 goods in vehicles used to carry passengers for hire or reward
- 52. Carriage of class 1 goods by road in motor vehicles
- 53. Marshalling and formation of trains
- 54. Keeping of information by carriers
- 55. Placards, marks and plate markings for carriage within Great Britain

PART 6

MISCELLANEOUS

- 56. Fees for applications relating to pressure receptacles and tanks
- 57. Transitional defence
- 58. Defence and enforcement
- 59. Amendments to the Health and Safety (Fees) Regulations 2004
- 60. Amendments to Chemicals (Hazard Information and Packaging for Supply) Regulations 2002
- 61. Amendments to the Classification and Labelling of Explosives Regulations 1983
- 62. Consequential amendments
- 63. Revocations and savings Signature

SCHEDULE 1 — OLD TANKS

- 1. Interpretation
- 2. Construction of tanks of old tank-vehicles, tank-containers and old tank wagons for carriage by road or rail
- 3. Testing, examination and maintenance for carriage by road or rail
- 4. Testing, examination and maintenance for carriage by road
- 5. Testing, examination and maintenance for carriage by rail
- 6. Use of old tank-vehicles, old tank wagons or tank-containers for carriage by road or rail
- 7. Keeping of documents
- 8. Appointment of inspection bodies by the competent authority
- 9. Exceptions

SCHEDULE 2 — OLD PRESSURE RECEPTACLES

- 1. Interpretation
- 2. Duties on those designing, manufacturing, importing, supplying, modifying or repairing old pressure receptacles
- 3. Conformity to approved design standard or specification
- 4. Examination of old pressure receptacles by competent or approved persons
- 5. Filling of old pressure receptacles
- 6. Approved design specification
- 7. Modification, repair and re-rating of old pressure receptacles
- 8. Additional requirements for old pressure receptacles containing certain dangerous goods not classified as class 2
- 9. Approvals by the competent authority
- 10. Exceptions

SCHEDULE 3 — COMPETENT AUTHORITY FUNCTIONS PART 1 — APPROVALS BY THE GB COMPETENT AUTHORITY

- 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...

- 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 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 other member States...
- 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 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...

- 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 8 — CONFORMITY MARKING

SCHEDULE 9 — PLACARDS, MARKS AND PLATE MARKINGS FOR CARRIAGE WITHIN GREAT BRITAIN

PART 1 — CARRIAGE OF GOODS BY ROAD

- 1. Where orange-coloured plates bearing a HIN are required to be...
- 2. Subject to paragraphs 3 and 6, where a transport unit...
- 3. Subject to paragraphs 4 and 6, where more than one...
- 4. Subject to paragraph 6, where more than one dangerous good...
- 5. (1) Subject to sub-paragraph (2), where dangerous goods are being...
- 6. (1) The information required to be displayed on placards and...

Hazard warning panel

PART 2 — CARRIAGE OF GOODS BY RAIL

- 7. Where orange-coloured plates bearing a HIN are required to be...
- 8. Where dangerous goods are being carried in tanks, a telephone...
- 9. (1) The information required to be displayed on placards and...

SCHEDULE 10 — AMENDMENTS TO THE HEALTH AND SAFETY (FEES) REGULATIONS 2004

- 1. The Health and Safety (Fees) Regulations 2004 shall be amended...
- 2. For regulations 12 to 14 substitute—Fees for certificates and...
- 3. For Schedule 10 substitute— SCHEDULE 10 FEE FOR DRIVER TRAINING...
- For Schedule 11 substitute— SCHEDULE 11 FEES FOR APPLICATIONS FOR
- 5. For Schedule 12 substitute— SCHEDULE 12 FEE FOR SAFETY ADVISER...

SCHEDULE 11 — AMENDMENTS TO THE CHEMICALS (HAZARD INFORMATION AND PACKAGING FOR SUPPLY) REGULATIONS 2002

- 1. The Chemicals (Hazard Information and Packaging for Supply) Regulations 2002...
- 2. After regulation 8 (labelling of dangerous substances and dangerous preparations)...

SCHEDULE 12 — AMENDMENTS TO THE CLASSIFICATION AND LABELLING OF EXPLOSIVES REGULATIONS 1983

- 1. The Classification and Labelling of Explosives Regulations 1983 shall be...
- 2. In regulation 2(1) (interpretation)—(1) omit the definition of "class...
- 3. In regulation 3 (classification and labelling of explosive articles and...
- 4. For regulation 6 (labelling of an article, substance or combination)...
- 5. Regulations 7 (labelling of an article, substance or combination not...

Status: This is the original version (as it was originally made).

- 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