
STATUTORY RULES OF NORTHERN IRELAND

2006 No. 173

HEALTH AND SAFETY

The Carriage of Dangerous Goods and Use of Transportable Pressure Equipment Regulations (Northern Ireland) 2006

Made - - - - *31st March 2006*

Coming into force *1st August 2006*

**THE CARRIAGE OF DANGEROUS GOODS AND
USE OF TRANSPORTABLE PRESSURE EQUIPMENT
REGULATIONS (NORTHERN IRELAND) 2006**

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Status: This is the original version (as it was originally made). This item of legislation is currently only available in its original format.

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- 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 Northern Ireland competent authority
- 9. Exceptions

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- 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
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- 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 Northern Ireland competent authority
- 10. Exceptions

SCHEDULE COMPETENT AUTHORITY FUNCTIONS

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PART I — APPROVALS BY THE NORTHERN IRELAND COMPETENT AUTHORITY

Status: This is the original version (as it was originally made). This item of legislation is currently only available in its original format.

1. The references referred to in regulation 27(b) are—
2. Any approval granted by the Northern Ireland competent authority by...
PART II — MULTILATERAL AND UNILATERAL APPROVALS BY THE NORTHERN IRELAND 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 Northern Ireland competent authority grants approvals under...
PART III — APPOINTMENT OF PERSONS BY THE NORTHERN IRELAND COMPETENT AUTHORITY
6. The references referred to in regulation 29(1)(b) are—
PART IV — RECOGNITION OF APPROVALS, TESTS, METHODS, STANDARDS, PROCEDURES ETC. BY THE NORTHERN IRELAND COMPETENT AUTHORITY
7. The references referred to in regulation 30(1)(b) are—
8. Where the Northern Ireland competent authority recognises a quality assurance...
PART V — IMPOSING OF REQUIREMENTS BY THE NORTHERN IRELAND COMPETENT AUTHORITY
9. The references referred to in regulation 31(1)(b) are—
10. (1) Where the Northern Ireland competent authority imposes requirements by...

SCHEDULE CONFORMITY ASSESSMENT PROCEDURES

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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 IV...
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

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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 IV...
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 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 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 ten 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...

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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 ten 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 ten 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 ten 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 pressure equipment***
 - 4.1 Each item of transportable pressure equipment must be individually examined...

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- 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 ten 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 ASSESSMENT
- SCHEDULE 6 CONFORMITY REASSESSMENT PROCEDURE
- 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
- 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...

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- 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 NORTHERN IRELAND
 - PART I — 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...
 - PART II — 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 REASONS FOR EXAMINATION NOT TAKING PLACE OR NOT BEING COMPLETED
 - 1. The applicant for the ADR certificate does not, after being...
 - 2. The particulars relating to the vehicle and shown in any...
 - 3. The vehicle is one as respects which it has been...
 - 4. The vehicle is a trailer, and is not accompanied by...
 - 5. There is not permanently affixed to the chassis or main...
 - 6. The vehicle or any motor vehicle by which it is...
 - 7. An inspector is not able to complete the inspection without...
 - 8. In the case of a trailer, an inspector is not...
 - 9. The vehicle or any trailer by which it is accompanied...
 - 10. An inspector is not able to complete the inspection due...
- SCHEDULE 11 AMENDMENTS TO THE CHEMICALS (HAZARD INFORMATION AND PACKAGING FOR SUPPLY) REGULATIONS (NORTHERN IRELAND) 2002
 - 1. The Chemicals (Hazard Information and Packaging for Supply) Regulations (Northern...

2. (1) In regulation 2(1), the definition of “the CDGCPL Regulations”...
- SCHEDULE 12 AMENDMENTS TO THE HEALTH AND SAFETY (FEES) REGULATIONS (NORTHERN IRELAND) 2005
1. The Health and Safety (Fees) Regulations (Northern Ireland) 2005 shall...
 2. For regulations 11 to 13 there shall be substituted the...
 3. For Schedule 8 there shall be substituted the following schedule—...
 4. For Schedule 9 there shall be substituted the following schedule—...
 5. For Schedule 10 there shall be substituted the following schedule—...
- SCHEDULE 13 CONSEQUENTIAL AMENDMENTS
1. Amendments to the Petroleum (Consolidation) Act (Northern Ireland) 1929
 2. Amendments to the Dangerous Substances in Harbour Areas Regulations (Northern Ireland) 1991
 3. Amendment to the Dangerous Substances (Notification and Marking of Sites) Regulations (Northern Ireland) 1992
 4. Amendment to the Notification of New Substances Regulations (Northern Ireland) 1994
 5. Amendment to the Health and Safety (Safety Signs and Signals) Regulations (Northern Ireland) 1996
 6. Amendments to the Reporting of Injuries, Diseases and Dangerous Occurrences Regulations (Northern Ireland) 1997
 7. Amendments to the Radiation (Emergency Preparedness and Public Information) Regulations (Northern Ireland) 2001
 8. Amendments to the Control of Asbestos at Work Regulations (Northern Ireland) 2003
 9. Amendment to the Control of Substances Hazardous to Health Regulations (Northern Ireland) 2003
 10. Amendment to the Control of Lead at Work Regulations (Northern Ireland) 2003
 11. Amendment to the Dangerous Substances and Explosive Atmospheres Regulations (Northern Ireland) 2003
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- Explanatory Note