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

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Coming into force 1st August 2006

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

PART I

INTRODUCTORY PROVISIONS

PART II

REQUIREMENTS OF ADR AND RID

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 4 CONFORMITY ASSESSMENT PROCEDURES

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

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...

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...