Directive 97/23/EC of the European Parliament and of the Council of 29 May 1997 on the approximation of the laws of the Member States concerning pressure equipment (repealed)

ANNEX I

ESSENTIAL SAFETY REQUIREMENTS

ANNEX II

CONFORMITY ASSESSMENT TABLES

1. . . . .

2. . . . .

3. . . . .

4. . . . .

ANNEX III

CONFORMITY ASSESSMENT PROCEDURES

. . . .

Module A (internal production control)
1. . . . .
2. . . . .
3. . . . .
4. . . . .
5. . . . .

Module A1 (internal manufacturing checks with monitoring of the final...

Module B (EC type-examination)
1. . . . .
2. . . . .
3. . . . .
4. . . . .
   - 4.1. . . . .
   - 4.2. . . . .
   - 4.3. . . . .
   - 4.4. . . . .
5. . . . .
6. . . . .
7. . . . .
8. . . . .
9. . . . .

Module B1 (EC design-examination)
1. . . . .
2. . . . .
3. . . . .
4. . . . .
   - 4.1. . . . .
   - 4.2. . . . .
   - 4.3. . . . .
5. . . . .
6. . . . .
7. . . . .

8. . . . .
9. . . . .

Module C1 (conformity to type)

1. . . . .
2. . . . .
3. . . . .
4. . . . .

Module D (production quality assurance)

1. . . . .
2. . . . .
3. Quality system
   - 3.1. . . . .
   - 3.2. . . . .
   - 3.3. . . . .
   - 3.4. . . . .
4. Surveillance under the responsibility of the notified body
   - 4.1. . . . .
   - 4.2. . . . .
   - 4.3. . . . .
   - 4.4. . . . .
5. . . . .
6. . . . .

Module D1 (production quality assurance)

1. . . . .
2. The manufacturer must draw up the technical documentation described below....
3. . . . .
4. Quality system
   - 4.1. . . . .
   - 4.2. . . . .
   - 4.3. . . . .
   - 4.4. . . . .
5. Surveillance under the responsibility of the notified body
   - 5.1. . . . .
   - 5.2. . . . .
   - 5.3. . . . .
   - 5.4. . . . .
6. . . . .
7. . . . .

Module E (product quality assurance)

1. . . . .
2. . . . .
3. Quality system
   - 3.1. . . . .
   - 3.2. . . . .
   - 3.3. . . . .
   - 3.4. . . . .
4. Surveillance under the responsibility of the notified body
   - 4.1. . . . .

4.2. . . . .
4.3. . . . .
4.4. . . . .
5. . . . .
6. . . . .

Module E1 (product quality assurance)
1. . . . .
2. The manufacturer must draw up the technical documentation described below....
3. . . . .
4. Quality system
4.1. . . . .
4.2. . . . .
4.3. . . . .
4.4. . . . .
5. Surveillance under the responsibility of the notified body
5.1. . . . .
5.2. . . . .
5.3. . . . .
5.4. . . . .
6. . . . .
7. . . . .

Module F (product verification)
1. . . . .
2. . . . .
3. . . . .
4. Verification by examination and testing of each item of pressure...
4.1. . . . .
4.2. . . . .
4.3. . . . .

Module G (EC unit verification)
1. . . . .
2. . . . .
3. . . . .
4. . . . .
4.1. . . . .
4.2. . . . .

Module H (full quality assurance)
1. . . . .
2. . . . .
3. Quality system
3.1. . . . .
3.2. . . . .
3.3. . . . .
3.4. . . . .
4. Surveillance under the responsibility of the notified body
4.1. . . . .
4.2. . . . .
4.3. . . . .

4.4. . . . .
5. . . . .
6. . . . .

Module H1 (full quality assurance with design examination and special...
1. . . . .
2. . . . .

ANNEX IV

MINIMUM CRITERIA TO BE MET WHEN DESIGNATING THE NOTIFIED BODIES REFERRED TO IN ARTICLE 12 AND THE RECOGNIZED THIRD-PARTY ORGANIZATIONS REFERRED TO IN ARTICLE 13

1. . . . .

2. . . . .

3. . . . .

4. . . . .

5. . . . .

6. . . . .

7. . . . .

ANNEX V

CRITERIA TO BE MET WHEN AUTHORIZING USER INSPECTORATES REFERRED TO IN ARTICLE 14

1. . . . .

2. . . . .

3. . . . .

4. . . . .

5. . . . .

6. . . . .

7. . . . .

ANNEX VI
CE MARKING

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ANNEX VII
DECLARATION OF CONFORMITY

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