

Directive 2009/23/EC of the European Parliament and of the  
Council of 23 April 2009 on non-automatic weighing instruments  
(Codified version) (Text with EEA relevance) (repealed)

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## ANNEX I

### ESSENTIAL REQUIREMENTS

Preliminary observation

### METROLOGICAL REQUIREMENTS

1. Units of mass
2. Accuracy classes
  - 2.1. ....
  - 2.2. Scale intervals
    - 2.2.1. The actual scale interval (d) and the verification scale interval...
    - 2.2.2. For all instruments other than those with auxiliary indicating devices:...
    - 2.2.3. For instruments with auxiliary indicating devices the following conditions apply:...
3. Classification
  - 3.1. Instruments with one weighing range
  - 3.2. Instruments with multiple weighing ranges
  - 3.3. Multi-interval instruments
    - 3.3.1. Instruments with one weighing range may have several partial weighing...
    - 3.3.2. Each partial weighing range i of multi-interval instruments is defined...
    - 3.3.3. The partial weighing ranges are classified according to Table 2....
4. Accuracy
  - 4.1. ....
  - 4.2. ....
5. ....
6. ....
7. Influence quantities and time
  - 7.1. ....
  - 7.2. ....
  - 7.3. ....
  - 7.4. ....
  - 7.5. ....
  - 7.6. ....
- Design and construction
8. General requirements
  - 8.1. ....
  - 8.2. ....
  - 8.3. ....
  - 8.4. ....
  - 8.5. ....
  - 8.6. ....
9. Indication of weighing results and other weight values
10. Printing of weighing results and other weight values
11. Levelling

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- 12. Zeroing
- 13. Tare devices and preset tare devices
- 14. ....
- 15. Price labelling instruments

## ANNEX II

### CONFORMITY ASSESSMENT PROCEDURES

- 1. EC type-examination
  - 1.1. ....
  - 1.2. ....
  - 1.3. The notified body shall:
    - 1.3.1. ....
    - 1.3.2. ....
    - 1.3.3. ....
    - 1.3.4. ....
  - 1.4. ....
  - 1.5. ....
  - 1.6. ....
  - 1.7. ....
- 2. EC declaration of type conformity (guarantee of production quality)
  - 2.1. ....
  - 2.2. ....
  - 2.3. Quality system
    - 2.3.1. The manufacturer shall lodge an application for approval of his...
    - 2.3.2. The quality system shall ensure conformity of the instruments with...
    - 2.3.3. The notified body shall examine and evaluate the quality system...
    - 2.3.4. ....
    - 2.3.5. ....
  - 2.4. EC surveillance
    - 2.4.1. ....
    - 2.4.2. The manufacturer shall grant the notified body access for inspection...
    - 2.4.3. ....
- 3. EC verification
  - 3.1. ....
  - 3.2. ....
  - 3.3. ....
  - 3.4. ....
  - 3.5. Verification by checking and testing of each instrument
    - 3.5.1. ....
    - 3.5.2. ....
    - 3.5.3. ....
- 4. EC unit verification
  - 4.1. ....
  - 4.2. ....
  - 4.3. ....
  - 4.4. ....

- 5. Common provisions
  - 5.1. ....
  - 5.2. ....
    - 5.2.1. ....
    - 5.2.2. ....
    - 5.2.3. ....
    - 5.2.4. ....

ANNEX III

DESIGN TECHNICAL DOCUMENTATION

- .....
- .....
- .....

ANNEX IV

‘CE’ CONFORMITY MARKING AND INSCRIPTIONS

- 1. Instruments subject to the EC conformity assessment procedure
  - 1.1. These instruments must bear:
  - 1.2. ....
  - 1.3. ....
  - 1.4. ....
  - 1.5. ....
- 2. Other instruments
- 3. Restrictive use symbol specified in Article 13

ANNEX V

The minimum criteria to be applied by member states when designating bodies for the carrying-out of tasks pertaining to the procedures referred to in article 9

- 1. ....
- 2. ....
- 3. ....
- 4. ....
- 5. ....
  - .....

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ANNEX VI  
'CE' CONFORMITY MARKING

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ANNEX VII

PART A

PART B

ANNEX VIII

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