



European
Commission

Regulation 339/93 on checks for conformity with the rules on product safety in the case of products imported from third countries

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Legal Aspects linked to Internal Market

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Background

Implementation of Internal Market

- **Free movement of goods**
- **Abolition of internal border controls**
 - **Close involvement of customs in market-monitoring operations**
 - **High level of health and safety**

Purpose of 339/93/EEC

**= Accompanying instrument
with twofold purpose**

- Legal basis for Customs to
suspend the release of unsafe
products**
- Install comparable rules
throughout Member States**

Products covered

- **ALL products**
 - Imported from third countries
 - Intended for release for free circulation
 - Whether or not falling under New Approach Directives
- **Except: Existence of specific Community provisions (e.g. plant health and veterinary controls)**

Commission Decision 1993/583

= List of products specifically affected regarding missing documents or marking

- **Toys (CE marking!)**
- **Medicinal products / human use**
- **Veterinary medicinal products**
- **Foodstuffs**

Procedure

Customs authorities

- **Risk to health or safety and/or**
- **Lack of document/marketing**
 - **Suspension of release of products for free circulation**
 - **Notification of national market surveillance authority**

Scenario 1

Serious and immediate risk to health or safety identified

- **Prohibition of placing on the market by MSA**
- **Endorsement on invoice + accompanying documents**

„Dangerous product – release for free circulation not authorized“

Scenario 2

Non-compliance with applicable product safety rules

- MSA take „appropriate action“, if necessary prohibiting the placing on the market
- Endorsement on invoice + accompanying documents
„Product not in conformity – release for free circulation not authorized“

Scenario 3

**No risk to health or safety,
no breach of product safety
rules**

**→ Release for free circulation
(provided all other conditions
regarding the release for free
circulation have been met)**

Scenario 4

**Customs are not notified
of any action taken by MSA**

- **Time limit: 3 working days from
suspension of release**

→ Release for free circulation



Implementation

Member States

- Designate MSA to be notified by customs
- Inform Commission of designated MSA to be notified by customs

Implementation

MSA must be in a position to

- **Verify compliance with product safety rules**
- **Within sufficiently short period**
- **Regarding any product the release of which is suspended**



The End

Thank you for your attention!