

Importing into the EU - Council Regulation (EEC) No 1991/2006

Report on the presentation held at BioFach, 17.02.2007, by Herman Van Boxem (European Commission, Agriculture and rural development Directorate-General Unit F5 - Organic farming)
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IMPORTS before 1.1.2007

All imports are based on an assessment of equivalency with EU Regulation 2092/91. There are two options:

- 1.) Third Country List
- 2.) Member State Authorizations

IMPORTS after 1.1.2007

Imports either have to meet EU Regulation 2092/91 (compliance) or have to meet equivalent standards to the EU Regulation. There are following options:

- 1.) Compliance
 - a.) control body list for compliance (not yet published, implementation procedures still need to be elaborated)
- 2.) Equivalence
 - a) Third Country List
 - b) Control Body List (not yet published, implementation procedures still need to be elaborated)
 - c) Member State Authorizations

EU Regulation 94/92 (Third Country List) and 1788/2001 (certificate of inspection for imports) remain in place and are linked to the new version of article 11.

EU Regulation 2092/91, Article 11:

- **Direct access: compliance**

11.1 Compliance

- a. product complies with articles 5 and 6 of 2092/91 (production rules)
- b. operators have submitted activity to inspection body/authority listed on new list (see 11.2)
- c. operators able to provide documentary evidence

11.2 Controls on compliance in third countries

- List of bodies/authorities by Commission
- Bodies ISO 65 accredited
- Bodies/authorities to send information
- Assessment reports by accreditation bodies or authorities
- Supervision by Commission and Member States

- **Equivalence**
 - 11.3 General conditions equivalence**
 - a. equivalent production standards
 - b. equivalent control arrangements
 - c. controls
 - by control system and control body on third country list – 11.4
 - by body on control body list – 11.5
 - d. product comes with certificate from bodies in c. (as in 1788/2001)
 - 11.4 List of third countries**
(As previously in 11.1), implementing rule (new 94/92)
 - Examination documented request third country
 - On-the-spot expert examination possible
 - Annual report
 - Commission and Member State supervision
 - 11.5 List of control bodies**
New (but elements from 11.6 of 2092/91)
Implementing rule: “second chapter to new 94/92”
 - Examination documented request control body
 - Regular on-the-spot evaluation
 - On-the-spot examination possible
 - Supervision by Commission and Member States based on assessment reports
 - SCOF decision
 - 11.6 Transition: prolongation of MS authorisations**
 - Prolongation of existing system of MS authorisations (article 11.6 of 2092/91) until 12 months after publication of first list of control bodies (new 11.5)
 - New: authorisation by Member State were the importer has notified his activity.
 - 11.7 Legal basis for implementing Commission rules**
 - a. For listing third countries and control bodies : “new 94/92”
 - b. For documentary evidence for compliant products (11.1.c)

Equivalence

- **Definition**

Capability to meet the same objectives and principles by applying rules which ensure the same level of assurance of conformity.

- **Reference**

Rules equivalent to EC rules. Assessment shall take into account Codex Alimentarius Guidelines CAC/GL 32