



# Import - Implementation rules, reasons and concerns

**Seminar at Biofach 2008**

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# Reasons for new import regulation

- > Action plan for organic food and farming of June 2004 (Action point 18): disadvantages of the current system – not accessible enough for operators in third countries; import certificate system not efficient
- > Deadline was in place to develop a better import system (End of 2006)
- > Now a new import regulation from 1. Jan 2007 has been decided – **what are the implications?**
- > **What were the main steps in this decision process?**

# Revision of the EU Regulation 2092/91 – what happened since last Biofach

- > **First draft December 2005 for a new Council regulation**
  - > Introduction of new system, including proposals for import
- > **From January 2006 until June 2007**
  - > intensive discussion process between the private sector, EU Member states and EU commission and partly Council and EU parliament
- > **Final version of the Council regulation (EC) No 1991/2006 on import 21 of December 2006**
  - > Import rules are already concluded and apply from 1 January 2007.
- > **Final Council regulation (EC) 834/2007 has been decided**
  - > Import rules are integrated in this regulation. Work on implementing rules started.

# Revision of the EU Regulation 2092/91 – what you should know?

- > Detailed EU commission implementing rules are in elaboration (put in force 1. Jan. 2009):
- > In January 2008 the EU Comissions made 2 proposals, one for production and one for import – currently discussed, **decided Mai/June 2009 (?)**
- > **What is the difference to old regulation EEC 2092/91:**
  - > Most of the annexes will be transposed without change - however with changed structure
  - > „Normal“ and „exceptional“ production rules – possibility for a certain regional flexibility (given criteria)
  - > New logo – mandatory in Europe – option for import.
  - > Indication of origin (EU agriculture / Non-EU agriculture )
  - > Import rules. *See presentation Herman van Boxem*



# Concerns and questions of the private sector, connected with the new import rules?

(e.g. from workshops of ORGAP Project, IFOAM EU commentary)

## General concerns:

- > **There will be 3 relevant documents, which are complementary – but this might be confusing for producers**
- > **There will be more flexibility for national governments within a given frame – but this needs transparent decision procedure**

## Regarding import

- > **The supervision of all proposed systems needs significant resources and capacities!**
- > **Avoidance of unnecessary burdens (allowance of electronic certificates)**
- > **Work of IFOAM-IOAS private accreditation system and Task force for harmonisation (FAO-OECD-IFOAM) should be taken into account**
- > **Guidelines are necessary for: accreditation of control bodies, equivalence assessment**