



1. New OCR _ EURLs/NRLs

2. WG microcriteria: *Salmonella* issues

22nd workshop of the EURL *Salmonella*
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Zaandam

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The new Official Control Regulation (EU) 2017/625 (OCR)

- *Published in the Official Journal on 7 April 2017*
- *Replaces Regulation (EC) No 882/2004 (and R 854/2004)*
- *Framework legislation on all controls by competent authorities including official sampling and analyses and requirements for official laboratories, NRLS and EURL*
- *To be completed by 34 delegated acts and 51 implementing acts*



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Structure of the OCR

General Principles (Articles 1 – 15)

Scope; definitions; transparency; responsibilities of Competent Authorities and business operators

Sector Specific Requirements (Articles 16 – 27)

e.g. products of animal origin; residues; animal welfare; GMOs; organic production; PDOs, PGIs, TSGs; new risks

Article 28–
33:
Delegation
of tasks

Article 34–
42:
Sampling,
analyses,
tests &
Diagnoses

Article 43–
76:
Import
controls

Article 77–
91:
Financing &
official
certification

Article 92–
101:
EURLs &
EURCs

Article 102–
108:
Administrative
Assistance &
Cooperation

Procedural and transitional provisions
(articles 142 – 167)



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Scope of the OCR (Art. 1.2)

Food and
food
safety

Feed and
feed
safety

GMOs

Animal
health

Animal
welfare

Animal
by-
products

Plant
health

Plant
protection
products

Organic
production

PDOs,
PGIs,
TSGs

Provisions on EURLs and NRLs

- *Title III (Art. 92-101) - Reference laboratories*
- *Key principles:*
 - **Broader scope (Plant Health)**
 - **Transparency and efficiency**
 - Decision and designation of EURLs
 - More specific and more precise requirements, responsibilities and tasks
 - **Accountability**
 - Distribution of responsibilities among COM, MS, EURLs and NRLs

Building-up the legislative framework

- ***Step 1- Decision to establish a EURL***
 - official controls depend on the quality, uniformity and reliability of methods and results
 - need to promote uniform practices in relation to the development or use of the methods
- ***Step 2- Designation of EURL***
 - Public selection process
 - Limited in time, with minimum period of 5 years or reviewed regularly
 - Accredited according to ISO/IEC 17025 (operating, methods of laboratory analysis and others and in a flexible manner)



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EURLs Requirements



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- **Staff:**
 - Impartial, confidentiality
 - Suitably, training + support
 - International Standards
 - Updated
- **Infrastructure, equipment + products**
 - According to needs
 - Emergency situations



- Equipped to comply with relevant biosecurity standards



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Responsibilities and tasks of EURLs

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- Improvement + harmonization of tests
- develop annual/multi-annual WP
- Scientific + technical assistance COM + NRLs (incl. training courses)
- assist during outbreaks



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- publish list of NRLs
- provide reference materials (if in WP)
- more specific:
 - proficiency tests, follow-up & inform COM + MS
 - cooperate to develop new methods
 - collaborate with Third Countries, with EFSA, EMA and ECDC



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Designation of NRLs

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- MSs to designate ≥ 1 NRLs for each EURL
- MS may designate lab in another MS/ 3rd Country
- NRL may be designated ≥ 1 MS



- MS may designate a NRL even if no EURL
- Official communication to COM, EURL + MS and public information of the name and address₉



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NRLs Requirements

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- Staff:
 - Impartial, confidentiality
 - Suitably and training + support
 - International Standards
- Infrastructure, equipment and products
 - According to needs
 - Emergency situations



- Scope of accreditation
- Equipped with relevant biosecurity standards
- CA shall organise audits
- Withdraw the designation if doesn't comply with:
 - ISO 17025
 - Obligations
 - Expected results in proficiency tests



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Responsibilities and tasks of NRLs



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- Methods of analysis + tests to official labs (OL)
- ensure dissemination to CA + OLs of info provided by EURL
- Scientific + technical assistance to CA for Multi Annual National Control Plan (MANCP)



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- Inform CA proficiency tests results and follow-up
- Conduct training courses for OLs
- Reference materials*
 - Validate
 - establish + maintain updated lists reference materials + manufacturers
- Assist CA in outbreaks



Obligations of Official Laboratories (OL)

Title II, Chapter IV, Article 38

- Inform immediately CA if results
 - indicate risk to human, animal or plant health, or as regards GMOs and plant protection products, also to the environment
 - or point to the likelihood of non-compliance
- Upon request by:
 - EURL or NRL ⇒ OL shall take part in proficiency tests.
 - Competent Authorities:
 - OL shall make available to the public methods used for analyses performed official controls
 - OL shall indicate the results together with the method used for each analysis



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Obligations of the MS

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- designate NRLs
- ensure coordination between NRLs work closely together
- communicate details of NRLs to COM, EURLs and other MS



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- Update and make available to public details NRLs (name and address)



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Timeline for EURLs

