

Official Controls Regulation 2017/625 – Overview and main changes

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I Legislative acts

REGULATIONS

* Regulation (EU) 2017/625 of the European Parliament and of the Council of 15 March 2017 on official controls and other official activities performed to ensure the application of food and feed law, rules on animal health and welfare, plant health and plant protection products, amending Regulations (EC) No 999/2001, (EC) No 396/2005, (EC) No 1069/2009, (EC) No 1107/2009, (EU) No 1151/2012, (EU) No 652/2014, (EU) 2016/429 and (EU) 2016/2031 of the European Parliament and of the Council, Council Regulations (EC) No 1/2005 and (EC) No 1099/2009 and Council Directives 98/58/EC, 1999/74/EC, 2007/43/EC, 2008/119/EC and 2008/120/EC, and repealing Regulations (EC) No 854/2004 and (EC) No 882/2004 of the European Parliament and of the Council, Council Directives 89/608/EEC, 89/662/EEC, 90/425/EEC, 91/496/EEC, 96/23/EC, 96/93/EC and 97/78/EC and Council Decision 92/438/EEC (Official Controls Regulation) (¹)

Scope of the OCR

Food and food safety

Feed and feed safety

GMOs

Animal health

Animal welfare

Animal byproducts

Plant health

Plant protection products

Organics

PDOs, PGIs, TSGs

Overview of the OCR

General Principles Articles 1 – 15

Subject matter, scope & definitions, Competent Authorities general requirements

Sector Specific Requirements Articles 16 – 27

e.g. Products of animal origin, residues, animal welfare, plant health, GMOs, plant protection products, organic production, new risks

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Art. 137–141 Enforce-ment

Common Provisions - Articles 142 – 167

Other official activities

The OCR applies to:

- Official controls (OC)
 - Tasks performed to verify compliance with the agri-food chain rules
- Other official activities (OOA)
 - Tasks carried out with the purpose to eliminate, contain or reduce any hazard which may arise for human, animal or plant health, animal welfare and the environment
 - Specific rules in the OCR apply also to OOA
 - Examples of OOA
 - Epidemiological surveillance and monitoring
 - Eradication and containment of diseases or pests
 - Issuance of official certificates or attestations

General rules

- Regular, unannounced risk-based controls to identify fraudulent or deceptive practices
- When planning controls to take account of:
 - operator's past record of compliance
 - the reliability of the operator's own checks
 - the likelihood of consumers being misled about nature, composition, etc. of food
- OC performed in a way that burden to operations is kept to minimum
- Financial penalties for fraudulent or deceptive practices would reflect the economic advantage or a percentage of the operator's turnover

General rules

- The OCR provides for CA to publish the outcome of OC regarding individual operators
 - Operators would be consulted first taking into account any comments made
- CA may publish information about the ratings of individual operators
- Increased transparency on calculation of fees and on breakdown of costs. Stakeholder consultation on method used to calculate fees
- Provisions for the protection of whistle blowers

Obligations of operators

- Provide details concerning name, legal form and specific activities they carry out
- Give access to computers, premises, documents, animals and goods under their control to the extent necessary to perform official controls or other official activities
- Assist and cooperate with the staff of the CA

Sampling, analyses, tests and diagnoses

- Rules on methods of sampling, analysis, tests and diagnoses apply to OC and OOA in all sectors covered by the OCR
- Operators' right to a second expert opinion at their own expense
- Provisions for official controls on animals and goods through distant communication
 - CA are not obliged to reveal their identity before they are in the possession of the samples.

Official laboratories (OL)

- OL may be designated in other Member States or third countries subject to certain conditions
- Participate to comparative or proficiency tests
- Make available to the public the methods used for analyses, tests or diagnoses and the results when requested by the CA
- OL to be regularly audited by CA
- Accreditation to EN ISO/IEC 17025
 - Temporary and permanent derogations

Import controls

- High risk animals and goods
 - Animals, POAO, germinal products, ABP, composite products, hay and straw, plants, plant products and other objects and certain feed and food of non-animal origin
 - Subject to controls at border control posts
 - 100% Doc checks, risk based identity and physical checks
 - Mandatory pre-notification requirement
 - Use of the Common Health Entry Document (CHED)

Import controls

- Low risk
 - Animals and goods which do not require systematic border controls
 - Controls performed at an appropriate place within the customs territory of the Union
 - Regular, risk based controls with appropriate frequency
 - Means of transport
 - Packaging (WPM)

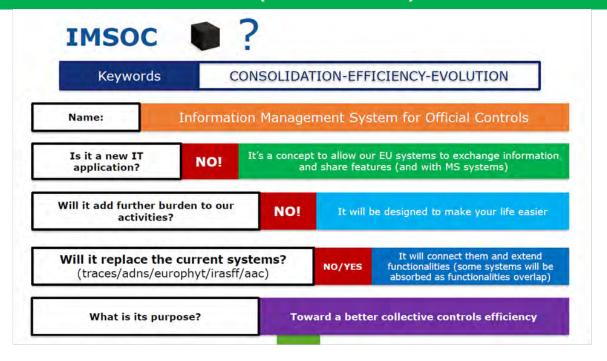
Import controls

- Physical checks performed on:
 - Plants, plant products and other objects by an official plant health officer
 - Animals (except aquatic animals), meat and edible meat offal by an official veterinarian (OV) assisted by trained staff
 - Aquatic animals, goods other than the above by OV or trained staff

Border Control Posts (BCPs)

- Designated facilities currently used for import controls, such as BIPs and DPEs will be replaced by BCPs
- They can be re-designated as BCP before the 14th December provided they meet the minimum requirements set out in the OCR and implementing regulation 2019/1014
- Additional conditions for animals, POAO, germinal products etc.

Information Management System for Official Controls (IMSOC)



New system from 14th — December

- TRACES Classic: exports, LMS, intra, DOCOMs
- TRACES-NT: imports
- Food and Feed Safety alerts (i-RASFF), AAC
- Animal Disease Information System (ADIS)
- EUROPHYT

IMSOC is an umbrella term to signify the electronic connection between the EU IT systems together to allow for better data exchange

CVED/CEDs → CHEDs

- TRACES-New Technology (NT) is a new system that will be used for imports from 14/12/19 and will host the Common Health Entry Documents (CHED). Developed for implementation of electronic-certification.
- A 'Common Health Entry Document' (CHED) will be used for all types of consignments entering the EU and subject to controls at BCPs
- There will be different CHEDs for different types of consignment e.g. CHED-D for high-risk FNAO will replace Common Entry Document (CED)
- IMSOC indicates to officials when to perform full checks → Risk-based controls, guided by IMSOC's predictive analysis



2 CHED rule on TRACES-NT

- Non-compliant consignments can be split into two: one for rejected and one for accepted (2 CHEDs required from original CHED)
- Second CHED is needed for:
 - Onward transportation
 - Inland Control Points

Electronic certification on TRACES-NT for imports into the EU

Paperless flow of SPS documentsofficials at BCPs can sign CHEDs electronically via e-signatures. (Some third Countries can sign EHCs electronically too). There are 2 ways to create an esignature:

- 1. Advanced: Password and one time password generated by SMS code
- Qualified: Password and one time password generated by token



- ❖ To set up → EC Trust Provider (LuxTrust) will contact individuals to set up esignatures
- Remember! There is always an option to wet sign paper CHEDs/EHCs

E-signatures

CHED should meet all of the following requirements:

- (i) it is signed with the electronic signature of the operator responsible for the consignment;
- (ii) it is signed with the advanced or qualified electronic signature of the certifying officer at border control posts or control points;
- (iii) it bears the advanced or qualified electronic seal of the issuing competent authority;
- (iv) it is sealed by TRACES with an advanced or qualified electronic seal;
- (v) each be timestamped with a qualified electronic time stamp

TRACES

- In principle, the OCR requires that from 14th December
 TRACES-NT must be used for all CHEDs
- TRACES Classic remains available for consultation purposes (+ INTRA, LMS, DOCOM, EXPORT)
 - No new CVEDs can be started (or replaced/modified)
 - New / In Progress CVEDs will change automatically to <u>Cancelled</u> status and must be re-done in TRACES.NT. For such cases, the Cancelled CVED can be consulted as reference
 - Control (Part III) can still be added to a CVED in a final status (arrival of goods, destruction...)