# Catalogue of Infringements Republic of Ireland

**Version 10** 



A catalogue of non-compliances, infringements and irregularities applying to the Organic Sector in Ireland, based on the provisions of the various EU regulations as well as national legislation.

Organic Unit, Department of Agriculture, Food and the Marine, Johnstown Castle Estate, Wexford, Ireland – January 2024

Updated: 1/12/2023

#### Introduction

This document is produced in accordance with Article 41, 42 and 43 of Regulation 848/2018<sup>1</sup> and Regulation 279/2021<sup>2</sup>.

Its purpose is to set out a comprehensive list of non-compliances, infringements and irregularities applying to the Organic Sector in Ireland, based on the provisions of the various EU regulations as well as national legislation.

#### Regulatory background

Article 41(4) of Regulation 848/2018 as amended provides as follows:

"Competent authorities shall provide a common catalogue of measures for cases of suspected non-compliance and established non-compliance to be applied in their territory, including by control authorities and control bodies".

#### Response of the Irish Competent Authority, the Department of Agriculture, Food and the Marine (DAFM)

The Irish Authorities hereby communicate to all control bodies a document outlining examples of types of non-compliance together with sanctions which should be imposed, at a minimum, and a catalogue listing infringements and irregularities affecting the organic status of products and corresponding measures to be applied by control bodies in cases of infringements or irregularities by operators under their control who are involved in organic production. In exceptional circumstances where the organic integrity of a product is directly compromised and after due consideration of mitigating circumstances, and with agreement of the Competent Authority, the sanction may be reduced. Similarly, the measures to be applied by the Organic Control Body (OCB) may be elevated to a higher sanction in proportion to the extent to which the provision has been violated, the type and circumstances of the irregularity and mindful of any pattern of reoccurrence. If the organic integrity of the product is not directly compromised then re-categorisation by the OCB is permitted. A composite list of all cases where sanctions are reduced must be maintained and available to the Department, as the Competent Authority, on request.

This document has been drawn up in consultation with The Organic Forum, representing all Control Bodies operating within the Irish jurisdiction. This document is not exhaustive and will be subject to on-going amendment. Other infringements and irregularities which also affect the organic status of products but are not listed must also be duly considered by the Control Body.

<sup>&</sup>lt;sup>1</sup> https://eur-lex.europa.eu/eli/reg/2018/848/2018-06-14

<sup>&</sup>lt;sup>2</sup> https://eur-lex.europa.eu/eli/reg\_impl/2021/279/oj

### **Note on the Suspension of Operators**

Regulation 848/2018<sup>3</sup>outlines, the additional rules on actions in case of non-compliance (Art. 41), additional rules on measures on the event of non-compliance (Art. 42) and additional rules on the exchange of information (Art. 43). The methodology for conducting an official investigation into the operator are outlined in Commission Implementing Regulation (EU) 2021/279, Art 2<sup>4</sup>.

In Ireland either type of breach (severe infringement or an infringement with prolonged effect) is referred to as a 'manifest infringement'. The period during which an operator is prohibited from marketing organic products should be considered on a case-by-case basis. However, to ensure that Irish Control Bodies adopt a broadly consistent approach, the following framework should be applied

The notion of "appropriate action" in the ISO 17065 Standard

Appropriate action can include:

- i. Continuation of the certification under conditions specified by the certification body (i.e., increased surveillance)
- ii. Reduction in scope of certification to remove non-conformity product variants
- iii. Suspension of the certification pending remedial action by the client
- iv. Withdrawal of the certification

#### The Framework

• Two years will generally be seen as the appropriate period of prohibition for a severe infringement or an infringement with prolonged effect. This prohibits the licensee to trade in any organic product for a period of two years. This should be regarded as the baseline against which other prohibitions are considered. A two-year period will in most cases allow sufficient time for operators to review their systems and implement compliant procedures. It should also offer organic consumers reassurance about the integrity of the organic sector.

<sup>&</sup>lt;sup>3</sup> https://eur-lex.europa.eu/eli/reg/2018/848/2018-06-14

<sup>&</sup>lt;sup>4</sup> https://eur-lex.europa.eu/eli/reg\_impl/2021/279/oj

The Tables in the Annexes provides more detail on:

- > The classifications of different types of non-compliance.
- > The types of actions that fall under each classification.
- > The action that Control Bodies are expected to take in respect of each type of classification.
- > The timescale for taking action; and
- any follow-up action that might be necessary.
- The following are examples of severe infringements or infringements with a prolonged effect:
  - > Fraudulent activity e.g., passing non-organic products off as organic.
  - > Incomplete records as a result of the omission of information.
  - > Two or more examples of behaviour that have a direct impact on the health and welfare of an operator's organic livestock assessed over a 12-month rolling period (depending on the severity of the case, one successful prosecution on these grounds could be sufficient to constitute a severe infringement or an infringement with prolonged effect); and
  - Failure, within a reasonable period, to correct three or more identified critical non-compliances.
- In determining the length of any prohibition, Control Bodies should consider both the circumstances of the breach and the circumstances of the operator. The following are examples of what are considered to be 'aggravating factors' and, if present alongside the severe infringement or infringement with prolonged effect, are likely to increase the prohibition period; this is not an exclusive list:
  - Evidence that animals under the case/control of the operator have been subjected to avoidable physical harm/mutilation/malnutrition that is inconsistent with the standard of care that is expected from an operator.
  - > The operator being obstructive towards any investigations undertaken by the Organic Control Bodies and/ or the Competent Authority following their findings.
  - > Actions that have resulted in a public health issue.
  - > Contamination of product due to inadequate measures to ensure separation of organic and non-organic products.
  - > Operator is unable to demonstrate the organic status of an ingredient used in a product.

In such cases, it may be considered appropriate to extend the agreed prohibition period to more than two years. This will be dependent on the individual circumstances of the matter.

DAFM and the Control Body should consider the individual case and agree a suitable period of prohibition. In exceptional circumstances after due consideration of mitigating circumstances, and with agreement of the Competent Authority, the sanction may be reduced.

• The Control Body should assess the circumstances surrounding the case and inform DAFM, suggesting a suitable period of prohibition. DAFM will then consider the case and confirm (with reasons) within 10 working days whether or not the suggested period of prohibition is considered to be appropriate. DAFM will ensure that all Control Bodies adopt a similar approach by checking as part of its annual assessments of the Control Bodies. Any non-compliances relating to the Control Body's additional private standards are not relevant under Article 42 Council Regulation (EC) 848/2018.

#### **End of the prohibition period:**

• Once the prohibition period ends, the operator can market products as organic provided they have a current organic licence, are registered with the Competent Authority and, where necessary, have complied with any requirements.

#### **Residue Analysis Action Form**

Any positive laboratory analysis result must be investigated by the OCB in order to determine the possible source of contamination and the appropriate follow-up action taken as outlined below.

The irregularity must be investigated in accordance with Regulation 848/2018 (Art. 28, 29)<sup>5</sup> and Commission Implementing Regulation (EU) 2021/279 (Art. 2)<sup>6</sup>, irregularities are notified to the Commission via the Organic Farming Information System (OFIS)

Residues	Status of contaminated	OCB Action Required	DAFM Action
	produce		
Below MRL	Temporarily hold affected organic batch for sale or distribution as organic for duration of investigation.	Investigate to identify possible source of contamination and make decision of action relation to product based on outcome of investigation.  Notify DAFM of residue finding, including investigation report, substantiated evidence and completed 'Notification of Irregularities' form.	Place notification on OFIS where necessary to notify other Member States or for an investigation to be carried out in a third country.
Residues detected greater than MRL	Immediately withdraw organic status of the product/lot/batch and prohibit its sale/distribution as organic. Instruct operator to quarantine product, notify customers of issue. Instruction from DAFM may require product recall and disposal.	Immediate notification to DAFM. Investigate to try to identify source of contamination. Based on outcome of investigation produce a report on same with findings and recommendations which may include suspension of operator and forward to DAFM. Send completed investigation report, substantiated evidence and completed 'Notification of Irregularities' form to DAFM.	DAFM will consult with relevant CA: DAFM/FSAI/SFPA with regard to the level of residue and the public health implications of consuming the product. DAFM will consider the case and confirm appropriate corrective action (with reasons) within 5 working days of receipt of the OCB report on contamination. Place notification on OFIS where necessary to notify other Member States or for an investigation to be carried out in a third country.

NB: In instances, where there is no MRL for the residue detected, the OCB should immediately contact DAFM for guidance.

<sup>&</sup>lt;sup>5</sup> https://eur-lex.europa.eu/eli/reg/2018/848/2018-06-14

<sup>&</sup>lt;sup>6</sup> https://eur-lex.europa.eu/eli/reg\_impl/2021/279/oj

#### Questions to be answered as part of the official investigation into an irregularity, and which are required on OFIS:

The investigation conducted by the OCB should aim to find the origin of the irregularity and the results of the investigations should determine the status of the product.

- (a) What type of investigation took place: physical, documentary checks?
- (b) Outline all the operators involved in the supply chain and their respective Control Bodies.
- (c) Traceability of the product should be provided; please provide COI details were applicable.
- (d) The name, batch number, quantity, ownership, and physical location of the organic or in-conversion products concerned.
- (e) Details on samples:
  - i. At which stage of production, preparation, or distribution and where exactly the presence of non-authorised products or substances has been detected, in particular for plant production, whether the sample was taken pre-harvest or post-harvest.
  - ii. Please provide analysis report.
- (f) If sampling was not possible no product in stock or the product should be sold, outline the amount of product in store.
- (g) The quantity of product placed on the market/ on hold/ withdrawn from the market.
- (h) Whether the products concerned are still placed on the market as organic or in-conversion products or used in organic production.
- (i) Whether the product has been downgraded to conventional or destroyed.
- (j) The type, name, quantity and other relevant information of the present non-authorised products or substances.
- (k) Whether other operators in the supply chain are affected, please provide details.
- (I) The results of previous official investigations on the products and operators concerned
  - i. Where there any severe infringement or prolonged infringement?
  - ii. Was the operator's certificate suspended/withdrawn & for how long?
- (m) the integrity of organic and in-conversion products.
- (n) the source and the cause of the presence of non-authorised products or substances.

# **ANNEXES**

Annex I Levels of non-compliance, definitions and examples

Annex II Catalogue of Infringements

Annex III Actions, Sanctions & Timescales

Annex I

# Levels of non-compliance, definitions, and examples

LEVEL 1 –	LEVEL 2 –	LEVEL 3 –	LEVEL 4 –
MINOR NON-	INTERMEDIATE NON-	CRITICAL NON-	MANIFEST
COMPLIANCE	COMPLIANCE	COMPLIANCE	INFRINCEMENT
Does not directly	May compromise the	The integrity of the	A serious and chronic failure
compromise the	integrity of the product if	operation, product/batch or	of the system where the
integrity of the	not corrected or may result	lot has been directly	integrity of the organic
product but needs	from not correcting a	compromised or lost but	production has been lost.
correcting,	previous minor non-	can be recovered.	Examples:
considering the	compliance.	For example:	Deliberate fraudulent
requirements under			activities such as
Reg. 2021/279, Art. 1	And where there is a	By accidental	substitution of non-organic
	suspicion that the cause of	use/substitution/	ingredients, selling non-
	the presence of the non-	contamination by	organic (n.o.) as organic
	authorised products or	prohibited materials	Contamination by
	substances lies under the	Non-compliant labelling	prohibited materials
	control of the operator, the	Excessive number of Non-	through systems failure
	operator shall examine any	compliances	The repeated failure to
	possible cause for the	<ul> <li>Contamination with GMOs</li> </ul>	correct previously identified
	presence of non-authorised		non-compliances
	products or substances.	The operator must inform	Livestock health & welfare
	(Reg. 2021/279, Art 1(1b)).	the OCB and provide	seriously compromised
		information and	Deliberate use of GMOs
	The operator must inform	documentation about the	
	the OCB and provide	supplier, traceability, lab	The operator must inform
	information and	results, sampling details and	the OCB and provide
	documentation about the	other relevant	information and
	supplier, traceability, lab	documentation.	documentation about the
	results, sampling details and	(Reg. 2021/279, Art 1) and	supplier, traceability, lab
	other relevant	the OCB conducts and	results, sampling details and
	documentation.		

	(Reg. 2021/279, Art 1(2))	official investigation (Reg. 2021/279, Art. 2).	other relevant documentation. (Reg. 2021/279, Art 1 and the OCB conducts and official investigation (Reg. 2021/279, Art. 2).
Example of non-com	pliance:		
Presence of pesticide	es-plant production products other than the	ose listed in the Regulation	
	Intermediate, if there is	Major, if there is evidence	Manifest Infringement, if
	evidence that the organic	that:	there is evidence that the
	operator:	The self-control	organic operators:
	> Has in plan	system is defective	use unauthorized
	adequate pest	(no pest	pesticides /plant
	management	management	protection
	procedures (i.e.,	procedures)	products/substances
	mechanical &	The organic	intentionally
	physical methods)	operator does not	
	> Uses pesticides	document under	
	/plant protection	what circumstance	
	products/substances	uses pesticides	
	only when the	/plant protection	
	above methods	products/substances	
	were not sufficient	> The organic	
	and only if they are	operator sourced	
	listed in the Annex	the unauthorized	
	➤ The use of	pesticides /plant	
	unauthorized	protection	
	pesticides /plant	products/substances	
	protection	from an un-reliable	
	products/substances	supplier	
	was just <u>one</u>	(unconsciously)	

incident due to non- compliance with self-control procedures  The product affected did not reach the market	
---	--

# Annex II CATALOGUE OF INFRINGEMENTS

				Level 1	Leve	el 2	Lev	el 3	Leve	el 4
Infringemen t Number	Compliance Category	Specific Non-Compliance Issue	Regulatory Reference <sup>7</sup>	Minor Non- Compliance	Intermediate Non- Compliance	Sanction	Critical Non- Compliance	Sanction	Manifest Infringement	Sanction
ORGANIC	PLANT & LIVES	TOCK PRODUCTION								
	Separation of Organic and Non-Organic Livestock	Organic and non-organic animals of same species on the same licensed holding (yards and land) simultaneously.	848.II.II.1.3.4.4.5				✓			
		Simultaneous grazing of organic and non-organic animals (different species) on same parcels	848.II.II.1.4.2				✓			
		Simultaneous grazing of organic and non-organic animals (different species) on same parcel in the contravention of grazing agreements.	848.II.II.1.4.2		✓					
		Non-organic animals exceeding grazing limit of 180 days	848.II.II.1.4.2		✓					
	The Grazing of Non-organic Grassland	Organic animals grazing non- organic land	848.II.II.1.4.2				✓			
	Origin of Livestock	Purchase of non-organic breeding stock which did not comply with nulliparous rule (exception rare breeds)	848.II.II.1.3				✓			

\_

<sup>&</sup>lt;sup>7</sup> Format of Regulation Notation: Reg x.Annex x.Part x.Subpoint x, e.g., 848.II.II.1.3, or Reg. Article x, e.g. 848.26;  $\Delta$  = Amended by

				Level 1	Leve	el 2	Leve	el 3	Leve	el 4
Infringemen t Number	Compliance Category	Specific Non-Compliance Issue	Regulatory Reference <sup>7</sup>	Minor Non- Compliance	Intermediate Non- Compliance	Sanction	Critical Non- Compliance	Sanction	Manifest Infringement	Sanction
		Did not apply for and/or receive derogation prior to purchase of non-organic stock, where eligible for a derogation	848.II.II.1.3		✓					
		Non-organic stock of incorrect gender bought in	848.II.II.1.3		✓					
		Purchase of nulliparous breeding stock in excess of 10% rule for bovines, or 20% rule for ovines, without derogation.	848.II.II.1.3				✓			
		Purchase of non-organic animals without derogation (40% rule) or in excess of derogation (40%)	848.II.II.1.3				✓			
		Purchase of 3-day old non-organic chicks without obtaining derogation in advance of purchase	848.II.II.1.3		✓					
		Purchase of non-organic young poultry over 3 days of age	848.II.II.1.3.4				✓			
		Failure to consult database/Organic Hub for	848. 26	<b>√</b>						
		purchase of organic stock	848.II.II.1.3.4.4							
	General Welfare & Management Issues	Animals (Bovines) not tagged/both tags missing in excess of DAFM timelines	Conditionality requirement				✓			
		Tagging of Stock (Bovines) not compliant, e.g., one tag missing	Conditionality requirement	✓						
		Derogation not sought for animal mutilations	848.II.II.1.7.8		✓					
		Mutilations not carried out in accordance with the legislation (i.e., anaesthetic/analgesia not used)	848.II.II.1.7.9				✓			
		Withdrawal periods non-compliant	848.II.II.1.5.2				✓			

				Level 1	Leve	el 2	Leve	el 3	Leve	el 4
Infringemen t Number	Compliance Category	Specific Non-Compliance Issue	Regulatory Reference <sup>7</sup>	Minor Non- Compliance	Intermediate Non- Compliance	Sanction	Critical Non- Compliance	Sanction	Manifest Infringement	Sanction
	Animal Housing	Animal bedding not provided	848.II.II.1.9				✓			
	Issues	Non-compliant animal bedding materials used	848.II.II.1.9		✓					
		Animals housed on slats with no access to bedded area	848.II.II.1.9				✓			
		Bedding over slats - no solid bedded area	848.II.II.1.9				✓			
		Inadequate animal bedding provided (i.e. comfortable, clean, dry rest area not evident; loose litter not evident over mats in cubicles):	848.II.II.1.9		✓					
		Animal housing – less than 50%								
		floor area solid, inadequate space for number of animals housed, inadequate perching space, pop- holes.	848.II.II.1.6 464/2020.1 464/2020.3 464/2020.4 464/2020.5				✓			
	Feed Issues	Feeding of non-organic feed to herbivores	848.II.II.1.9				✓			
		Feeding non-organic feed to non- herbivores in excess of 5%	848.II.II.1.9.3 848.II.II.1.9.4				✓			
		Feeding non-organic feed containing GM	848.11						✓	
		Supplementary feeding of mineral licks containing GM material	848.11				✓			
		First year in-conversion fodder utilised in excess of 20% of overall fodder requirements –from the organic farm	848.II.II.1.4.3		✓					

				Level 1	Leve	el 2	Leve	el 3	Leve	el 4
Infringemen t Number	Compliance Category	Specific Non-Compliance Issue	Regulatory Reference <sup>7</sup>	Minor Non- Compliance	Intermediate Non- Compliance	Sanction	Critical Non- Compliance	Sanction	Manifest Infringement	Sanction
		Feeding in-conversion feed in excess of allowances stipulated in regulations	848.II.II.1.4.3				<b>✓</b>			
	Animal Health & Welfare Issues	Animals do not have access to open air areas when conditions allow or have access to roughage (except pigs, poultry, and bees)	848.II.II.1.4.1e				✓			
		Use of veterinary inputs without adequate justification	848.II.II.1.5.2		✓					
		Use of substances having hormonal or thryostatic action and beta agonists in farm animals without veterinary authorisation on a case-by-case basis.	848.II.II.1.5				<b>✓</b>			
		Evidence of inadequate provision of feed, water and other necessary substances which compromises animal health/welfare	848.6				✓			
		Specific nutritional requirements set out in the legislation of the animal have not been met	848.II.II.1.4 848.II.II.1.9.1.1 848.II.II.1.9.2.1 848.II.II.1.9.3.1 848.II.II.1.9.4.2 848.II.II.1.9.5.1 848.II.II.1.9.6.2				<b>✓</b>			
		Specific animal welfare requirements set out in the legislation have not been met	848.II.II.1.7 848.II.II.1.9.4.3 848.II.II.1.9.6.5				✓			
		Specific housing and husbandry requirements set out in the legislation of the animal have not been met	848.II.II.1.6 848.II.II.1.9.1.2 848.II.II.1.9.2.2 848.II.II.1.9.3.2 848.II.II.1.9.4.4 848.II.II.1.9.5.2				<b>✓</b>			

				Level 1	Leve	el 2	Leve	el 3	Leve	el 4
Infringemen t Number	Compliance Category	Specific Non-Compliance Issue	Regulatory Reference <sup>7</sup>	Minor Non- Compliance	Intermediate Non- Compliance	Sanction	Critical Non- Compliance	Sanction	Manifest Infringement	Sanction
		Specific health care requirements of the animals set out in the legislation have not been met	848.II.II.1.5 848.II.II.1.9.6.3				✓			
	Land and Pollution	Crop rotation without mandatory legumes not in compliance	848.II.II.1.9		✓					
	Related Issues	Soil analysis or justification for an input not available	848.II.I.1.9.3		✓					
		Severe poaching of soil	848.II.II.1.7 848.II.II.1.9				✓			
		Exceeding annual limit relating to 170 kgs/ON/ha/pa	848.II.I.1.9 848.II.II.1.6				✓			
		Manure storage/effluent storage and management non-compliant (e.g. storing manure on land during closed period)	848.II.I.1.9 848.II.II.1.6				✓			
	Livestock Paperwork	Inaccurate stock figures stock reconciliation not possible	848.39				✓			
	Issues	Flock/Herd register/ CMMS not up to date in accordance with statutory regulations.	848.II.II.1.3 Δ 2021/1691.34.8				✓			
		Livestock Sales Declaration form not available for organic stock purchased			✓					
		Veterinary Health Plan not up to date	848.II.II.1.5 Δ 2021/1691.34.8	✓						

				Level 1	Lev	el 2	Leve	el 3	Leve	el 4
Infringemen t Number	Compliance Category	Specific Non-Compliance Issue	Regulatory Reference <sup>7</sup>	Minor Non- Compliance	Intermediate Non- Compliance	Sanction	Critical Non- Compliance	Sanction	Manifest Infringement	Sanction
		No Veterinary Health Plan	848.II.II.1.5 Δ 2021/1691.34.8		✓					
		Proof of use of anaesthetic/analgesia not available					✓			
		Documentation not submitted by specified deadline	848.39		✓					
	General Paperwork Issues	No Records kept	848.39 848.II.I.1.12						<b>✓</b>	
		Inadequate record-keeping	848.39 848.II.I.1.12 Δ 2021/1691				✓			
		Extension/reduction of licensed land areas not notified to OCB	848.39		✓					
		Organic enterprise changes not notified to OCB, e.g. approval not sought for new enterprise and/or product	848. 39		✓					
		Discrepancy in mass balance audit; mass balance audit not achievable	848.39 848.II.I.1.12		✓					
	Seed Paperwork Issues	Derogation not sought for use of untreated non-organic seed or propagation material (100% n.o.)	848.II.I.1.8 Δ 2020/1794. 12.2b		<b>✓</b>					
		Derogation not sought for permission to use seed mixture containing a % n.o. seed	848.II.1.1.8 Δ 2020/1794. 12.2b		<b>✓</b>					

				Level 1	Leve	el 2	Leve	el 3	Leve	el 4
Infringemen t Number	Compliance Category	Specific Non-Compliance Issue	Regulatory Reference <sup>7</sup>	Minor Non- Compliance	Intermediate Non- Compliance	Sanction	Critical Non- Compliance	Sanction	Manifest Infringement	Sanction
		Use of chemically dressed/treated seeds	848.II.I.1.8 Δ 2020/1794. 12.2b				✓			
		Failure to consult database and Organic Hub for purchase on non-organic seed	848.26		✓					
		Marketing of organic or in- conversion plant reproductive material (heterogenous) without prior notification to competent authority	848.13		✓					
		Marketing of organic plant reproductive material that is not in compliance with the regulation	2021/1189.3				<b>✓</b>			
		Parallel Production in Crop Production	848.9				✓			
	Prohibited Inputs & Contamination	Spraying prohibited herbicide/pesticide	848.II.I.1.10 Δ 2021/1691.I.1b				✓			
	Risks	Failure to report a known spray- drift issue	848.39		✓					
		Use of prohibited chemical(s), inputs	848.II.I.1.10 Δ 2021/1691.I.1b				✓			
		Exceeding limit for copper usage Where copper product approved for use in Ireland as a fungicide	848.II.I.1.10 848.24 Δ 2021/1691.I.1b				✓			
		Cleanliness of equipment not in compliance	848.II.I.1.11 Δ 2021/1691.I.1c		✓					
		Cleaning procedures not adequately recorded (use, date, name, active substance and location of use)	848.34(8) 2021/1691.I.4(c)		✓					

				Level 1	Lev	el 2	Leve	el 3	Leve	el 4
Infringemen t Number	Compliance Category	Specific Non-Compliance Issue	Regulatory Reference <sup>7</sup>	Minor Non- Compliance	Intermediate Non- Compliance	Sanction	Critical Non- Compliance	Sanction	Manifest Infringement	Sanction
		Use of compost for propagation purposes which contains inputs other than those indicated in Regulations.	848.II.I.1.9 Δ 2021/1691.I.1a				<b>✓</b>			
		Storage of prohibited input on an organic holding	848. 24				✓			
		Inadequate Precautionary Measures in respect of contamination (Equipment, utensils, housing, pens etc.)	848.II.I.1.6	✓						
	General Issues	Export of organic manure/poultry litter/slurry to non-organic farms	848.II.I.1.9		✓					
		Persistent failure to correct previous issues of critical non-compliance	848.41 848.42 848.43							
			2017/625.138 279/ 2021.I							
ORGANIC PR	OCESSING & PRO	CESSED PRODUCTS								
		Point of sale labelling non- compliant (e.g. No compulsory indications whatsoever)	848.30 848.32				✓			
	Product or Labelling Issues	Some compulsory indications missing or incorrect on organic product packaging (e.g. EU logo)	848.32848.33		✓					
	133463	Display signage non-compliant: Loose products only – out of date certificate on display, activity not covered, product/product category not listed	848.30		✓					
		Use of unapproved non-organic ingredient in an organic product	848.30				<b>✓</b>			
		Sale of non-organic produce as 'organic'	848.30						✓	

				Level 1	Lev	el 2	Leve	el 3	Leve	el 4
Infringemen t Number	Compliance Category	Specific Non-Compliance Issue	Regulatory Reference <sup>7</sup>	Minor Non- Compliance	Intermediate Non- Compliance	Sanction	Critical Non- Compliance	Sanction	Manifest Infringement	Sanction
		Use of unapproved processing aid or additive	848.24				✓			
		Use of non-rinse sanitiser without subsequent rinsing	848.II.IV.1.5		✓					
		Use of unapproved off-site processing unit	848.34				✓			
		Use of non-food grade packaging on organic food products	848.2				✓			
		Segregation between organic/in- conversion/. non-organic products not evident/not compliant	848.II.IV.1.5				<b>✓</b>			
		Agreed bleed runs/purges not carried out between organic/inconversion/ non-organic production runs	848.II.IV.1.5				<b>✓</b>			
		Organic products in storage not identifiable	848.II.IV.1.5				✓			
		Clean-down prior to organic production run not evident/non-compliant	848.II.IV.1.5				✓			
		Use of unlicensed wholesaler or unlicensed storage facility	848.34				✓			
		Pest Infestation in food store not addressed	848.II.IV.1.5				✓			
		Insufficient action taken on complaints	848.27		✓					
		Feed production not in compliance with regulation (processor)	848.17848.III.V 848.III.2.1.2				✓			

				Level 1	Leve	el 2	Lev	el 3	Leve	el 4
Infringemen t Number	Compliance Category	Specific Non-Compliance Issue	Regulatory Reference <sup>7</sup>	Minor Non- Compliance	Intermediate Non- Compliance	Sanction	Critical Non- Compliance	Sanction	Manifest Infringement	Sanction
		Feed labelling not in compliance with the legislation (processor)	848.117 848.111.2.1.2848.3 2				<b>✓</b>			
		Some compulsory indications missing or incorrect on organic feed labelling	848.III.2.1.2 848.32		✓					
		Collection, packaging and storage of products not in compliance with the regulation	848.27 848.III.V				<b>✓</b>			
		Failure to keep records of feed, formulations	848.II.V Δ 2021/1691.I.5(c)				✓			
		Records not kept of location and quantity of wild plants collected	848.II.I. 2.2 Δ 2021/1691.I.1(e)				✓			
	Residue and Analysis Issues	Failure to notify OCB of positive residue test result taken as part of licensee's own analysis procedures	848.27				✓			
		DNA analysis reveals DNA other than DNA of specific product, e.g., pork DNA in beef burger produced by organic licensee	848.7				✓			
		Product has been irradiated (as evidenced by irradiation test) – in product produced by licensee	848.5(i)				✓			
	Product Paperwork	Documentation not submitted by specified deadline	2017/625.15		✓					
		Failure to notify CB immediately of any irregularity/infringement or suspicion that may impact on the organic status of a product	848.27				✓			

				Level 1	Leve	el 2	Leve	el 3	Leve	el 4
Infringemen t Number	Compliance Category	Specific Non-Compliance Issue	Regulatory Reference <sup>7</sup>	Minor Non- Compliance	Intermediate Non- Compliance	Sanction	Critical Non- Compliance	Sanction	Manifest Infringement	Sanction
		Inadequate record-keeping	848.39 as amended, delegated reg. of July 2021				✓			
		Proof of GM-free status of non- organic permitted ingredients not verifiable	848. 5f 848.III.11				✓			
		Flavourings in use not compliant with regulatory requirements	848.24 848.16				✓			
		Pre-approval not sought for production of new products	848.39				✓			
		Processing records - quantities of ingredients etc not adequate to production	848.39 as amended, delegated reg. of July 2021				<b>✓</b>			
		Purchase invoices not stating organic status of ingredient being brought in	848.II.I. 1.12 848.39 as amended, delegated reg. of July 2021		✓					
		Sales invoice/docket not stating organic status of product	848.39 as amended, delegated reg. of July 2021	✓						
		Insufficient/inadequate records to complete mass balance	848.II.I.1.12 848 Article 39 as amended, delegated reg. of July 2021				<b>✓</b>			
			848.II.I.1.12							
		Input/Output does not balance –	2021/279.9				<b>√</b>			
		over usage Traceability not achievable due to inadequate record-keeping	848.38(8) 848.38(8)				<b>✓</b>			

				Level 1	Leve	el 2	Leve	el 3	Leve	el 4
Infringemen t Number	Compliance Category	Specific Non-Compliance Issue	Regulatory Reference <sup>7</sup>	Minor Non- Compliance	Intermediate Non- Compliance	Sanction	Critical Non- Compliance	Sanction	Manifest Infringement	Sanction
		Pest control records inadequate/non-compliant	848.39 as amended, delegated reg. of July 2021 848.II.I.1.12		✓					
		Poor hygiene standards in evidence in production unit	848.II.IV (Processed food production rules)				✓			
		Requirements for Third country imports Certificate of Inspection not met	2021/2307 and 2021/2325				✓			
		Import of organic and inconversion products from 3 <sup>rd</sup> Countries not in compliance with the regulation	848.45				✓			
		Specifications for ingredients not available	848.II.V.1.1 848II.IV.1.3				✓			
		Incoming organic goods not checked on arrival/inadequate verification of goods received	848.111.5				✓			
		Delivery dockets for bulk products non-compliant as regards the required organic certification ID	848.39 as amended, delegated reg. of July 2021				<b>✓</b>			
		Persistent failure to correct previous issues of critical non-compliance	848.II.I.1.12 848.41 848.42 848.43							
			2017/ 625.138 (Actions in the event of established non- compliance)						✓	
			279/ 2021.I							

				Level 1	Leve	el 2	Leve	el 3	Leve	el 4
Infringemen t Number	Compliance Category	Specific Non-Compliance Issue	Regulatory Reference <sup>7</sup>	Minor Non- Compliance	Intermediate Non- Compliance	Sanction	Critical Non- Compliance	Sanction	Manifest Infringement	Sanction
		No Organic product recall system	848.27							
			848.41				<b>V</b>			
		Organic Product recall conducted without notifying OCB	848.27				✓			
			848.41							
ORGANIC A	QUACULTURE & AQ	UACULTURE PRODUCTS								
	General Issues	Disease prevention/veterinary treatments non-compliant	848.11.111.3.1.4				✓			
		Withdrawal periods non- compliant	848.11.111.3.1.4				✓			
		Transport of live fish not in compliance with regulations	848.11.111.1.7				<b>✓</b>			
		Sustainability Management Plans not updated	848.11.111.1.5		✓					
		Inadequate record-keeping	848.39				✓			
		Measures taken against predators	848.II.I.1.12							
		not recorded in Sustainable  Management Plan	848.11.111.3.2.2		✓					
		Origin of aquatic animals not in compliance with regulations	848.II.III.3.1.2 and Δ 2020/1693						✓	
		Slaughtering techniques and/ or handling prior to slaughter non-compliant.	848.II.III.3.1.6				✓			
		On land rearing system non- compliant	848.II.III.3.1.5				✓			
		Use of hormone or hormone derivatives	848.11.111.3.1.2				✓			
		Feed not in compliance with Regulations	848.II.II.3.1.3							
			848.II.III.3.1.2 Δ				✓			
			2020/427.I.1.3a							

				Level 1	Lev	el 2	Leve	el 3	Leve	el 4
Infringemen t Number	Compliance Category	Specific Non-Compliance Issue	Regulatory Reference <sup>7</sup>	Minor Non- Compliance	Intermediate Non- Compliance	Sanction	Critical Non- Compliance	Sanction	Manifest Infringement	Sanction
		Antibiotic Residues following analysis where use of antibiotic was not prescribed by vet	848.II.I.1.7	848.II.II. 1.5.2			✓			
		Persistent failure to correct previous issues of critical non-compliance	848.41 848.42 848.43							
			2017/ 625.138 (Actions in the event of established non- compliance)						✓	
		Failure to consult database and Organic Hub for purchase on non- organic aquaculture juveniles	279/2021.I 848.26		✓					
		Aquaculture production - Failure to keep records (origin of animals, feeding regimes and disease prevention measures)	848.II.III Δ 2021/1691.I.III				✓			
		Inadequate separation of organic and non-organic production units	848.II.I.2.1						✓	
		Use of veterinary medicines not declared to Control Bodies.	848.11.111.3.1.4				<b>✓</b>			
	Mollusc Production	Simultaneous production not in compliance with Regulations	848.9				✓			
		Production area not delineated as required	848.11.111.3.2.2				✓			
		Seed not sourced in compliance with Regulations	848.11.1.1.8				✓			

				Level 1	Lev	el 2	Lev	el 3	Leve	el 4
Infringemen t Number	Compliance Category	Specific Non-Compliance Issue	Regulatory Reference <sup>7</sup>	Minor Non- Compliance	Intermediate Non- Compliance	Sanction	Critical Non- Compliance	Sanction	Manifest Infringement	Sanction
		Harvesting of organic mussels in Class B Waters that do not have High Ecological status	848.II.III.3.1.3.2				<b>✓</b>			
	Finfish Production	Maximum stocking densities exceeded	848.15 848.3 848.4				✓			
		Full description of production site not available	848.39		<b>✓</b>					
	Seaweed Production	Cleaning and drying of seaweed not compliant	848.11.111.2				<b>✓</b>			
		Bio-fouling organisms removed by means other than physical	848.11.111.3.1.4				<b>✓</b>			

# **Annex III**

# **Actions, Sanctions and Timescales**

NB Examples within the relevant definition's sections are not exhaustive and will be subject to on-going additions/amendments

Level	Description	Definition	Action / Sanctions	Time Scale	Follow-up
0	Compliance	Fully compliant.  No issues raised.	None.	N/A	None.
0	Comment or Observation	The means of notifying general information regarding the standards.  Example – references to:  Practices that could be improved e.g. to best practise. Interpretation of the standards laid down in the organic Regulations. Forthcoming changes to the standards.	<ul> <li>Request for information /clarifications</li> <li>Notice of non-compliance/Order to correct non-compliance(s) within an agreed timeframe</li> </ul>	N/A	Must be checked at subsequent inspection
1	MINOR non-compliance	Does not directly compromise the integrity of the product but needs correcting	<ul> <li>Corrective action to be agreed in writing by the CB and operator.</li> <li>Operator to commit to undertake corrective action within an agreed timetable.</li> <li>Evidence of compliance to be supplied by operator and verified by CB.</li> <li>Only where evidence of compliance cannot be supplied, a statement of intent may be accepted (e.g. where a long term capital investment is required).</li> </ul>	Licensee to respond within time period set by the CB, not exceeding 30 days from the date of notification.  Corrective actions to be implemented within a reasonable period agreed by the CB taking account of the type of non-compliance (e.g. whether just a minor technical matter (such as	Must be checked at subsequent inspection

			<ul> <li>Reduction in scope: to exclude products or activities which do not meet the certification requirements</li> <li>The operator can invoke an appeals procedure</li> <li>Notice of non-compliance/Order to correct non-compliance(s) within an agreed timeframe</li> <li>Provisional prohibition of placing the product on the market</li> </ul>	record keeping) or potentially having wider repercussions (e.g. on livestock welfare) if not corrected.	
2	INTERMEDIATE non-compliance	May compromise the integrity of the product if not corrected or may result from not correcting a previous minor non-compliance.	<ul> <li>Corrective action to be agreed in writing by the CB and operator.</li> <li>Operator to commit to undertake corrective action within an agreed timetable.</li> <li>Evidence of compliance to be supplied by operator and verified by the CB.</li> <li>Only where evidence of compliance cannot be supplied a statement of intent may be accepted (e.g. where a long term capital investment is required).</li> <li>Suspension: during the suspension period the organic operator is not allowed to market any products as inconversion or organic</li> <li>The operator can invoke an appeals procedure</li> <li>Decertification of the parcel (s) or the entire farm to lower conversion stage (i.e. from year 2 to year 1) or conventional status</li> <li>Decertification of products from inconversion and organic status to conventional</li> </ul>	Licensee to respond within time period set by the CB, not exceeding 30 days from the date of notification.  Corrective actions to be implemented within a reasonable period agreed by the CB taking account of the type of non-compliance.	An additional inspection may be required, at the discretion of the CB.  Corrective actions to be verified at subsequent inspection.

			Renewal of certification under conditions     Reduction in the scope of the certification     Suspension of the certification     Product recall		
3	CRITICAL non-compliance	The integrity of the operation, product/batch or lot has been directly compromised or lost but can be recovered – Examples:  By accidental use/substitution/contamination with prohibited materials.  Non-compliant labelling.  Excessive number of non-compliances.	The regulatory requirement here is to ensure that product affected (production run or entire lot) is not marketed as organic (having due regard to principle of proportionality). The EU regulations also require immediate notification to other OCBs, Competent Authority and relevant Member States, as well as EU Commission if appropriate (Reg. 848/2018 and Implementing Reg. 2021/279)  Withdrawal: the operator can no longer market any products as in-conversion or organic  termination of certification agreement  The operator can invoke an appeals procedure  Withdrawal of the Certification	Decertification of land, product, batch, lot as appropriate with immediate effect.	Before the suspension can be lifted:  The operator provides evidence that the critical noncompliance has been corrected.  Additional inspection at the discretion of the CB to check for full compliance (e.g. only where the suspension was found to be justified).  Corrective action and status of decertified land, product, batch, lot to be checked at subsequent inspection.
4	MANIFEST INFRINGEMENT Severe infringements and	A serious and chronic failure of the system where the integrity of the organic production has been lost.  Examples:	The regulatory requirement here is to ensure that product affected (production run or entire lot) is not marketed as organic (having due regard to principle of proportionality). The EU regulations also require immediate notification to other OCBs, Competent	To be agreed between DAFM and the OCB.	The Control Body and DAFM to agree on a period during which the

infringements with prolonged effect	<ul> <li>Deliberate fraudulent activities such as substitution of non-organic ingredients, marketing non organic produce as organic.</li> <li>Contamination by prohibited materials through systems failure.</li> <li>The repeated failure to correct previously identified non-compliances.</li> </ul>	Authority and relevant Member States, as well as EU Commission if appropriate. (EU Reg 834/2007 Arts 30.1 & 30.2)  Immediate notification to DAFM Immediate verbal suspension/ decertification.  Referred to emergency meeting of the CB's Certification Committee for confirmation/decisions. The Certification Committee meeting may be teleconference or email.  Decertification confirmed in writing by	The oper apply for licence fr	rganic products. rator may not ran organic rom another d OCB during od of
		<ul> <li>days, but no more than seven working days.</li> <li>DAFM informed of decision &amp; CBs if product recall is needed.</li> <li>FSAI to be notified by DAFM as appropriate.</li> <li>Period of licence withdrawal to be agreed with DAFM</li> <li>DAFM to notify other Member States and Commission as required.</li> </ul>		

#### Precautionary measures to avoid the presence of non-authorised products and substances (Re. 848/2018, Article 28)

- 1. In order to avoid contamination with products or substances that are not authorised in accordance with the first subparagraph of Article 9(3) for use in organic production, operators shall take the following precautionary measures at every stage of production, preparation and distribution:
- (a) put in place and maintain measures that are proportionate and appropriate to identify the risks of contamination of organic production and products with non-authorised products or substances, including systematic identification of critical procedural steps;
- (b) put in place and maintain measures that are proportionate and appropriate to avoid risks of contamination of organic production and products with non-authorised products or substances;
- (c) regularly review and adjust such measures; and
- (d) comply with other relevant requirements of this Regulation that ensure the separation of organic, in-conversion and non-organic products.
- 2. Where an operator suspects, due to the presence of a product or substance that is not authorised pursuant to the first subparagraph of Article 9(3) for use in organic production in a product that is intended to be used or marketed as an organic or in-conversion product, that the latter product does not comply with this Regulation, the operator shall:
- (a) identify and separate the product concerned;
- (b) check whether the suspicion can be substantiated;
- (c) not place the product concerned on the market as an organic or in-conversion product and not use it in organic production unless the suspicion can be eliminated;
- (d) where the suspicion has been substantiated or where it cannot be eliminated, immediately inform the relevant competent authority, or, where appropriate, the relevant control authority or control body, and provide it with available elements, where appropriate;
- (e) fully cooperate with the relevant competent authority, or, where appropriate, with the relevant control authority or control body, in identifying and verifying the reasons for the presence of non-authorised products or substances.