ANIMAL HEALTH ADVISORY COMMITTEE
Working Group of the Advisory Group on the Food Chain

Brussels, 12 October 2021

Amendments to Commission Delegated Regulation (EU) 2020/687 on disease control measures

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Commission Delegated Regulation (EU) 2021/1140

- ✓ Article 21(3) has been clarified;
- ✓ Article 22 and Annex VI have been amended to clarify provisions as regards ABPs;
- ✓ Article 112 (‘Repeals’) has been replaced by the following:

> 1. Directives 92/66/EEC, 2001/89/EC, 2003/85/EC and 2005/94/EC as well as the acts adopted on the basis of those Directives, shall cease to apply with effect from 21 April 2021.
>
> 2. **Directives 2000/75/EC and 2002/60/EC as well as the acts adopted on the basis of those Directives, shall cease to apply with effect from 14 July 2021.’;**

Draft delegated act on the use of veterinary medicinal products for prevention and control of listed diseases

Supplementing Article 47(1) of the AHL

SANTE/7144/2020

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Legal basis (AHL)

Article 46(1)

Provides for the possibility for the Member States to take measures concerning **the use of veterinary medicinal products (ALL)** to ensure the most efficient **prevention or control** of listed diseases **(ALL)**. These measures may cover prohibitions, restrictions and compulsory use of veterinary medicinal products and must be previously assessed as proportionate and necessary.

Article 47(1)

Empowers the Commission to adopt delegated acts concerning:

- ✓ prohibitions and restrictions on the use of veterinary medicinal products;
- ✓ specific conditions for the use of veterinary medicinal products for a specific listed disease;
- ✓ risk-mitigation measures to prevent the spread of listed diseases through animals treated with the veterinary medicinal products or products from such animals;
- ✓ surveillance for specific listed diseases following the use of vaccines and other veterinary medicinal products.

Process

✓5 Expert group meetings (March 2020-September 2021)

✓Discussions focused on the scope as regards:

- Listed diseases
- Animal species
- Veterinary medicinal products

✓Based on these discussions, a first draft was presented, with a wide scope and including general rules for the use of VMPs

New approach

Rules on the use of **certain VMPs** for prevention and control of **certain listed diseases (Part I) – Terrestrial and aquatic animals**

Circumstances under which **vaccines for category A** diseases can be used

Which **VMPs cannot be used for category A and B** diseases

Rules on the use of **vaccines** for prevention and control of category A diseases **(Part II) – Terrestrial animals**

- Preconditions
- Strategies
- General rules

Disease-specific rules **(Annexes)**

- Implementation
- Measures (prohibitions) in the vaccination zone
- Recovery of the previous animal health status

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Next steps

- ✓Next Expert Group meetings in October and November
  - ✓Discussion focused on disease-specific rules
- ✓Submission for public consultation
- ✓Adoption: 2Q 2022

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Amendments to Commission Delegated Regulation (EU) 2020/692 on entry into the Union

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Commission Delegated Regulation (EU) 2021/1705

- ✓ Correction of mistakes, clarifications, new possibilities not provided for in the original text following discussions with MSs, TC and stakeholders
- ✓ Amendments to:
  - ✓ Articles 1, 2, 3, 13, 26, 49, 57, 60, 62, 74, 80, 83, 85, 87, 91, 100, 102, 107, 110, 111, 119, 125, 154, 167, 169, 172, 173, 174, 175, 182
  - ✓ Annexes III, VIII, XV, XXVIII, XXIX

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Commission Delegated Regulation (EU) 2021/1703 (‘Composite products’)

- ✓ Article 162 has been amended to include ‘colostrum based products’.
- ✓ Article 163 has been amended to allow:
  - ✓ the entry into the Union of shelf-stable composite products containing dairy products that originate from third countries listed for the entry into the Union of raw milk and dairy products not subject to a risk-mitigating treatment, without undergoing any specific risk-mitigating treatment;
  - ✓ the entry into the Union of shelf-stable composite products containing dairy products that originate from third countries listed for the entry into the Union of dairy products subject to a risk-mitigating treatment, if they have undergone a risk-mitigating treatment in accordance with Article 157 of Delegated Regulation (EU) 2020/692.

Commission Delegated Regulation (EU) 2021/1703

Dairy products contained in shelf-stable composite products (not containing meat products)

No specific treatment	Treatments in column A of Annex XXVI to CDR (EU) 2020/692	Treatments in column B of Annex XXVI to CDR (EU) 2020/692
TC or zones listed in Annex XVII to CIR (EU) 2021/404	TC or zones listed in Annex XVIII to CIR (EU) 2021/404	TC or zones that are not listed in Annex XVII or XVIII to CIR (EU) 2021/404

But they have to be listed for entry of meat products, egg products or fishery products

These amendments will be reflected in point 10 of the model attestation provided for in Annex V to CIR (EU) 2020/2235

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Thank you

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