EUROPEAN UNION **Export Health Certificate**

Part I : Details of consignment

I.1. Consignor
Name
Address
Country ISO Code

I.2. IMSOC Reference
Specimen not to be used for exports from EU

I.2.a. Local Reference

I.5. Consignee
Name
Address
Country ISO Code

I.3. Central competent authority

I.4. Local competent authority

I.7. Country of origin ISO Code

I.9. Country of destination ISO Code

I.8. Region of origin Code

I.10. Region of destination Code

I.11. Place of Dispatch
Name
Address
Approval Number
Country ISO Code

I.12. Place of destination
Name
Address
Approval Number
Country ISO Code

I.13. Place of Loading
Name
Address
Approval Number
Country ISO Code

I.14. Date and time of departure

I.15. Means of Transport

Mode	International transport document	Identification

I.16 Entry Point

I.18. Transport conditions
Ambient ☐ Controlled temperature ☐ Frozen ☐ Chilled ☐

I.17. Accompanying documents
Commercial document reference Date of issue
Country Place of issue

I.19. Container No / Seal No

I.20. Certified as

Transhumance ☐	Pollination ☐	Animal Feedingstuff ☐	Pharmaceutical use ☐
Game Restocking ☐	Fattening ☐	Registered equidae ☐	Quarantine ☐
Artificial reproduction ☐	Training ☐	Further process ☐	Racing ☐
Production of petfood ☐	Breeding and production ☐	Consignments according to Regulation No 999/2001 ☐	Competition ☐
Storage ☐	Breeding ☐	Ornamental use/research ☐	Laboratory ☐
Rodent food ☐	Pets ☐	Other ☐	Production ☐
Technical use ☐	Ornamental bird food ☐	Relaying ☐	Slaughter ☐
Approved Bodies ☐	Human consumption ☐	Pet food ☐	Sales ☐
Organic fertilizers ☐	Unregistered equidae ☐	Circus exhibition ☐	Category 3 fish oil/fish meal for detoxification according to Regulation 2015/786 ☐

I.21. For transit through a third country ☐
Country ISO Code
EU Exit Authority BCP code
EU Entry Authority BCP code

I.22. For transit through Member State(s) ☐
Country ISO Code

I.23. Total number of packages	I.24. Total quantity	I.25. Total net weight	I.25. Total gross weight

Part I : Details of consignment

I.28. Description of consignment

1. 04 DAIRY PRODUCE; BIRDS' EGGS; NATURAL HONEY; EDIBLE PRODUCTS OF ANIMAL ORIGIN, NOT ELSEWHERE SPECIFIED OR INCLUDED

0401 Milk and cream, not concentrated nor containing added sugar or other sweetening matter

Commodity	Species	Quantity	Batch number	Manufacturing plant

Cold store	Cutting plant	Date of freezing	Date of production	Date of slaughter

Net weight	Product Description	Package count	Identification mark

Part II: Certification

II. Health information

II.1. Animal Health Attestation

I, the undersigned official veterinarian, declare that I am aware of the relevant provisions of Directive 2002/99/EC and of Regulation (EC) No 853/2004 and hereby certify that the ☐ colostrum/ ☐ colostrum-based products (1) described in Part I:

have been obtained or manufactured from colostrum obtained from animals:

(i) under the control of the official veterinary service;

(ii) which were in a third country or part thereof that has been free of foot-and-mouth disease and of rinderpest for a period of at least 12 months prior to the date of this certificate, and where vaccination against foot-and-mouth disease has not been carried out during that period;

(iii) belonging to holdings which were not under restrictions due to foot-and-mouth disease or rinderpest, and

(iv) subject to regular veterinary inspections to ensure that they satisfy the animal health conditions laid down in Chapter I of Section IX of Annex III to Regulation (EC) No 853/2004 and in Directive 2002/99/EC.

II.2. Public Health Attestation

I, the undersigned official inspector, declare that I am aware of the relevant provisions of Regulations (EC) No 178/2002, (EC) No 852/2004, (EC) No 853/2004 and (EC) No 854/2004 and hereby certify that the ☐ colostrum/ ☐ colostrum-based products made with colostrum(1) described in Part I were produced in accordance with those provisions, and in particular that:

(a) they were manufactured from colostrum:

(i) which comes from holdings registered in accordance with Regulation (EC) No 852/2004 and checked in accordance with Annex IV to Regulation (EC) No 854/2004;

(ii) which was produced, collected, cooled, stored and transported in accordance with the hygiene conditions laid down in Chapter I of Section IX of Annex III to Regulation (EC) No 853/2004;

(iii) which complies with the guarantees on the residues status of colostrum provided by the monitoring plans for the detection of residues or substances submitted in accordance with Directive 96/23/EC, and in particular, Article 29 thereof;

(iv) which, pursuant to testing for residues of antibacterial drugs carried out by the food business operator in accordance with the requirements of point 4 in Part III of Chapter I of Section IX of Annex III to Regulation (EC) No 853/2004, complies with the maximum residue limits for residues of antibacterial veterinary medicinal products laid down in the Annex to Regulation (EU) No 37/2010;

(v) which has been produced under conditions guaranteeing compliance with the maximum residue levels for pesticides laid down in Regulation (EC) No 396/2005, and maximum levels for contaminants laid down in Regulation (EC) No 1881/2006;

(b) they come from an establishment implementing a programme based on the HACCP principles in accordance with Regulation (EC) No 852/2004;

(c) they have been processed, stored, wrapped, packaged and labeled in accordance with Chapters III and IV of Section IX of Annex III to Regulation (EC) No 853/2004;

(d) they meet the relevant requirements laid down in Chapter II, Section IX of Annex III to Regulation (EC) N0 853/2004 and the relevant microbiological criteria laid down in Regulation (EC) No 2073/2005 on microbiological criteria for foodstuffs, and

(e) the guarantees covering live animals and products thereof provided by the residue plans submitted in accordance with Directive 96/23/EC, and in particular Article 29 thereof are fulfilled.

Part II: Certification

II. Health information

Notes

(*) Those countries subject to the transitional import arrangements include: an EU member State; Iceland; Liechtenstein; Norway and Switzerland.

References to Great Britain in this certificate include Channel Islands and Isle of Man.

References to European Union legislation within this certificate are references to direct EU legislation which has been retained in Great Britain (retained EU law as defined in the European Union (Withdrawal) Act 2018).

This certificate is intended for colostrum or colostrum—based products from third countries or parts thereof authorised in column A of Annex 1 to Regulation (EU) No 605/2010.

Part I:

Box reference I.7: Provide name and ISO code of the country or part thereof as appearing in Annex 1 to Commission Regulation (EU) No 605/2010 of 2 July 2010 laying down animal and public health and veterinary certification conditions for the introduction into the European Union of raw milk and dairy products intended for human consumption.

Box reference I.11: Name, address and approval number of the establishment of dispatch.

Box reference I.15: Registration number (railway wagons or container and road vehicle), flight number (aircraft) or name (ship). In case of unloading and reloading, the consignor must inform the border control post of introduction into Great Britain.

Box reference I.16: Do not use this box until the end of the transitional staging period.

Box reference I.19: Use the appropriate Harmonised System (HS) code under the following headings: 04.01; 04.02; 04.03; 04.04; 04.05; 04.06; 04.10; 15.17; 17.02; 19.01; 21.05; 21.06; 22.02; 28.35; 30.01; 35.01; 35.02 or 35.04.

Box reference I.20: Indicate total gross weight and total net weight.

Box reference I.23: For containers or boxes, the container number and the seal number (if applicable) should be included.

Box reference I.28: Manufacturing plant: introduce the approval number of the production holding(s), collection centre or standardization centre approved for exportation to Great Britain.

Part II:

(1) Keep as appropriate.

The colour of the signature shall be different to that of the printing. The same rule applies to stamps other than those embossed or watermark.

Certifying Officer

Name (in capital letters)	Qualification and title
Date of signature	Signature
Stamp	

Část I

I.1. Odesílatel
Název
Adresa
Země Kód ISO

I.2. Referenční číslo IMSOC
Specimen not to be used for exports from EU

I.2.a. Local Reference

I.5. Příjemce
Název
Adresa
Země Kód ISO

I.3. Ústřední příslušný orgán

I.4. Local competent authority

I.7. Země původu Kód ISO

I.9. Country of destination Kód ISO

I.8. Region of origin Kód

I.10. Region určení Kód

I.11. Place of Dispatch
Název
Adresa
Číslo schválení
Země Kód ISO

I.12. Místo určení
Název
Adresa
Číslo schválení
Země Kód ISO

I.13. Místo nakládky
Název
Adresa
Číslo schválení
Země Kód ISO

I.14. Date and time of departure

I.15. Dopravní prostředky

Typ	Doklad	Identifikace

I.16 Entry Point

I.18. Transport conditions
Okolní ☐ Controlled temperature ☐ Zmrazené ☐ Chlazený ☐

I.17. Průvodní doklady
Referenční číslo obchodního do k ladu
Datum vydání
Země
Místo vydání

I.19. Č. kontejneru / č. plomby

I.20. Certified as

Transhumance ☐	Pollination ☐	Krmivo pro zvířata ☐	Použití pro farmaceutické účely ☐
Game Restocking ☐	Výkrmová ☐	Evidovaní koňovití ☐	Karanténa ☐
Umělé rozmnožování ☐	Training ☐	Další postup ☐	Racing ☐
Production of petfood ☐	Breeding and production ☐	Consignments according to Regulation No 999/2001 ☐	Competition ☐
Storage ☐	Breeding ☐	Ornamental use/research ☐	Laboratory ☐
Rodent food ☐	Zvířata v zájmovém chovu ☐	Jiné ☐	Production ☐
Technické použití ☐	Ornamental bird food ☐	Sádkování ☐	Porážka ☐
Schválené orgány ☐	Lidská spotřeba ☐	Krmivo pro domácí zvířata ☐	Sales ☐
Organic fertilizers ☐	Unregistered equidae ☐	Cirkus/výstava ☐	Category 3 fish oil/fish meal for detoxification according to Regulation 2015/786 ☐

I.21. For transit through a third country ☐
Země Kód ISO
EU Exit Authority BCP code
EU Entry Authority BCP code

I.22. For transit through Member State(s) ☐
Země Kód ISO

I.23. Celkový počet balení	I.24. Celkové množství	I.25. Celková čistá hmotnost	I.25. Celková hrubá hmotnost

Část I

I.28. Description of consignment

1. 04 MLÉKO A MLÉČNÉ VÝROBKY; PTAČÍ VEJCE; PŘÍRODNÍ MED; JEDLÉ PRODUKTY ŽIVOČIŠNÉHO PŮVODU, JINDE NEUVEDENÉ ANI NEZAHRNUTÉ

0401 Mléko a smetana, nezahuštěné, neobsahující přidaný cukr ani jiná sladidla

Komodita	Druh	Množství	Číslo šarže	Výrobní zařízení

Chladírenské zařízení	Bourárna	Datum zmrazení	Datum výroby	Datum porážky

Čistá hmotnost	Product Description	Počet balení	Identifikační značka

Part II: Certification

II. Informace týkající se zdraví

II.1. Animal Health Attestation

I, the undersigned official veterinarian, declare that I am aware of the relevant provisions of Directive 2002/99/EC and of Regulation (EC) No 853/2004 and hereby certify that the ☐ colostrum/ ☐ colostrum-based products (1) described in Part I:

have been obtained or manufactured from colostrum obtained from animals:

(i) under the control of the official veterinary service;

(ii) which were in a third country or part thereof that has been free of foot-and-mouth disease and of rinderpest for a period of at least 12 months prior to the date of this certificate, and where vaccination against foot-and-mouth disease has not been carried out during that period;

(iii) belonging to holdings which were not under restrictions due to foot-and-mouth disease or rinderpest, and

(iv) subject to regular veterinary inspections to ensure that they satisfy the animal health conditions laid down in Chapter I of Section IX of Annex III to Regulation (EC) No 853/2004 and in Directive 2002/99/EC.

II.2. Public Health Attestation

I, the undersigned official inspector, declare that I am aware of the relevant provisions of Regulations (EC) No 178/2002, (EC) No 852/2004, (EC) No 853/2004 and (EC) No 854/2004 and hereby certify that the ☐ colostrum/ ☐ colostrum-based products made with colostrum(1) described in Part I were produced in accordance with those provisions, and in particular that:

(a) they were manufactured from colostrum:

(i) which comes from holdings registered in accordance with Regulation (EC) No 852/2004 and checked in accordance with Annex IV to Regulation (EC) No 854/2004;

(ii) which was produced, collected, cooled, stored and transported in accordance with the hygiene conditions laid down in Chapter I of Section IX of Annex III to Regulation (EC) No 853/2004;

(iii) which complies with the guarantees on the residues status of colostrum provided by the monitoring plans for the detection of residues or substances submitted in accordance with Directive 96/23/EC, and in particular, Article 29 thereof;

(iv) which, pursuant to testing for residues of antibacterial drugs carried out by the food business operator in accordance with the requirements of point 4 in Part III of Chapter I of Section IX of Annex III to Regulation (EC) No 853/2004, complies with the maximum residue limits for residues of antibacterial veterinary medicinal products laid down in the Annex to Regulation (EU) No 37/2010;

(v) which has been produced under conditions guaranteeing compliance with the maximum residue levels for pesticides laid down in Regulation (EC) No 396/2005, and maximum levels for contaminants laid down in Regulation (EC) No 1881/2006;

(b) they come from an establishment implementing a programme based on the HACCP principles in accordance with Regulation (EC) No 852/2004;

(c) they have been processed, stored, wrapped, packaged and labeled in accordance with Chapters III and IV of Section IX of Annex III to Regulation (EC) No 853/2004;

(d) they meet the relevant requirements laid down in Chapter II, Section IX of Annex III to Regulation (EC) N0 853/2004 and the relevant microbiological criteria laid down in Regulation (EC) No 2073/2005 on microbiological criteria for foodstuffs, and

(e) the guarantees covering live animals and products thereof provided by the residue plans submitted in accordance with Directive 96/23/EC, and in particular Article 29 thereof are fulfilled.

Part II: Certification

II. Informace týkající se zdraví

Notes

(*) Those countries subject to the transitional import arrangements include: an EU member State; Iceland; Liechtenstein; Norway and Switzerland.

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This certificate is intended for colostrum or colostrum—based products from third countries or parts thereof authorised in column A of Annex 1 to Regulation (EU) No 605/2010.

Part I:

Box reference I.7: Provide name and ISO code of the country or part thereof as appearing in Annex 1 to Commission Regulation (EU) No 605/2010 of 2 July 2010 laying down animal and public health and veterinary certification conditions for the introduction into the European Union of raw milk and dairy products intended for human consumption.

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Part II:

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Certifying Officer

Name (in capital letters) | Qualification and title

Datum podpisu | Podpis

Razítko