MODEL ANIMAL HEALTH CERTIFICATE FOR THE MOVEMENT BETWEEN MEMBER STATES OF OVINE AND CAPRINE ANIMALS NOT INTENDED FOR SLAUGHTER (MODEL 'OV/CAP-INTRA-X')

EUROPEAN UNION					INTRA
Part I: Description of consignment	I.1	**Consignor** Name Address Country ISO country code		I.2 **IMSOC reference** I.2a **Local reference** I.3 **Central Competent Authority** I.4 **Local Competent Authority**	**QR CODE**
	I.5	**Consignee** Name Address Country ISO country code		I.6 **Operator conducting assembly operations independently of an establishment** Name Registration No Address Country ISO country code	
	I.7	**Country of origin** ISO country code		I.9 **Country of destination** ISO country code	
	I.8	**Region of origin** Code		I.10 **Region of destination** Code	
	I.11	**Place of dispatch** Name Registration/Approval No Address Country ISO country code		I.12 **Place of destination** Name Registration/Approval No Address Country ISO country code	
	I.13	**Place of loading**		I.14 **Date and time of departure**	
	I.15	**Means of transport** □ Vessel □ Aircraft □ Railway □ Road vehicle Identification □ Other Document		I.16 **Transporter** Name Registration/Authorisation No Address Country ISO country code I.17 **Accompanying documents** Type Code Country ISO country code Commercial document reference	
	I.18	**Transport conditions** □ Ambient		□ Chilled □ Frozen	
	I.19	**Container number/Seal number** Container No Seal No			

I.20	**Certified as or for**		
□ Further keeping	□ Slaughter	□ Confined establishment	□ Germinal products
□ Registered equine animal	□ Travelling circus/animal act	□ Exhibition	□ Event or activity near borders
□ Release into the wild	□ Dispatch centre	□ Relaying area/purification centre	□ Ornamental aquaculture establishment
□ Further processing	□ Organic fertilizers and soil improvers	□ Technical use	□ Quarantine or similar establishment
□ Products for human consumption	□ Pollination	□ Live aquatic animals for human consumption	□ Other

I.21	**□ For transit through a third country**	
	Third country	ISO country code
	Exit point	BCP code
	Entry point	BCP code

I.22	**□ For transit through Member State(s)**		**I.23**	**□ For export**	
	Member State	ISO country code		Third country	ISO country code
	Member State	ISO country code		Exit point	BCP code
	Member State	ISO country code			

I.24	**Estimated journey time**	**I.25**	**Journey log**	□ yes	□ no
I.26	**Total number of packages**	**I.27**	**Total quantity**		
I.28	**Total net weight/gross weight (kg)**	**I.29**	**Total space foreseen for the consignment**		

I.30 Description of consignment

CN code	Species	Subspecies/Category	Sex	Identification system	Identification number	Age	Quantity Type
Region of origin		Cold store		Identification mark	Type of packaging		Net weight
Slaughterhouse		Treatment type		Nature of commodity	Number of packages		Batch No
		Date of collection/production		Manufacturing plant	Approval or registration number of plant/establishment/centre	Test	

Part II: Certification

II. Health information

II.a Certificate reference	II.b IMSOC reference

I, the undersigned official veterinarian, hereby certify that:

II.1. The ovine/caprine animals(1) of the consignment described in Part I meet the following requirements:

II.1.1. They are identified as provided for in Article 45(2) or (4) or Article 46(1) of Commission Delegated Regulation (EU) 2019/2035.

II.1.2. They, for at least the 30 day period prior to the departure of the consignment, or since birth, if they are younger than 30 days of age,

II.1.2.1. have been continuously resident in the establishment of origin;

II.1.2.2. have not been in contact with kept ovine or caprine animals of a lower health status or subject to movement restrictions for animal health reasons;

II.1.2.3. have not been in direct or indirect contact with kept animals that have entered the Union from a third country or territory during the 30 day period prior to the departure of the animals.

II.1.3. They have not shown clinical signs or symptoms of diseases listed for ovine/caprine animals during the clinical examination which was carried out, within the 24 hour period prior to departure of the consignment, on ……. (*insert date dd/mm/yyyy*).

II.2. According to official information, the animals described in Part I meet the following health requirements:

II.2.1. They do not come from establishments subject to movement restrictions affecting the species or situated in a restricted zone established for reasons of diseases listed for ovine/caprine animals.

(2)*either*[II.2.2. They come from establishments free from infection with *Brucella abortus, B. melitensis* and *B. suis* without vaccination regarding ovine and caprine animals, and

(2)*either* [the establishments of origin are situated in a Member State or zone thereof with the status free from infection with *Brucella abortus, B. melitensis* and *B. suis* regarding the ovine and caprine population;]

(2)*and/or* [they have been subjected to a test for infection with *Brucella abortus, B. melitensis* and *B. suis* with one of the diagnostic methods provided for in Part 1 of Annex I to Commission Delegated Regulation (EU) 2020/688, carried out, with negative results, on a sample taken during the 30 day period prior to departure, and in the case of post-parturient females taken at least 30 days after parturition;]

(2)*and/or* [they are less than 6 months old;]

(2)*and/or* [they are castrated.]

(2)*or*[II.2.2. They come from establishments free from infection with *Brucella abortus, B. melitensis* and *B. suis* with vaccination regarding ovine and caprine animals and they are moved to a Member State or zone thereof without the status free from infection with *Brucella abortus, B. melitensis* and *B. suis* regarding ovine and caprine animals.]

(2)*either*[II.2.3. They are kept ovine animals and come from establishments in which infection with *Mycobacterium tuberculosis* complex (*M. bovis, M. caprae and M. tuberculosis*) has not been reported during the last 42 days prior to departure.]

(2)*and/or*[II.2.3. They are kept caprine animals and come from establishments in which surveillance for infection with *Mycobacterium tuberculosis* complex *(M. bovis, M. caprae* and *M. tuberculosis)* has been carried out on the caprine animals kept on the establishments during at least the 12 month period prior to departure, as referred to in Article 15(3) of Delegated Regulation (EU) 2020/688.]

	II.2.4.	They come from establishments in which infection with rabies virus in kept terrestrial animals has not been reported during the 30 day period prior to departure.
	II.2.5.	They come from establishments situated in an area of at least 150 km radius around those establishments in which infection with epizootic haemorrhagic disease virus has not been reported in kept animals of listed species for that disease during the last 2 years prior to departure.
	II.2.6.	They come from establishments in which anthrax in ungulates has not been reported during the 15 day period prior to departure.
	II.2.7.	They come from establishments in which surra (*Trypanosoma evansi*) has not been reported during the 30 day period prior to departure, and
	(2)*either*	[surra has not been reported in the establishments during the last 2 years prior to their departure.]
	(2)*or*	[surra has been reported during the last 2 years prior to departure, following the last outbreak the affected establishments have remained under movement restrictions until: – the infected animals have been removed from the establishments, and – the remaining animals on the establishments have been subjected to a test for surra (*Trypanosoma evansi*) with one of the diagnostic methods provided for in part 3 of Annex I to Delegated Regulation (EU) 2020/688, carried out, with negative results, on samples taken at least 6 months after the infected animals have been removed from the establishments.]
	(2)[II.2.8.	They are kept uncastrated male ovine animals, and – come from establishments in which ovine epididymitis (*Brucella ovis*) has not been reported during the 12 month period prior to departure, and – have been subjected to a serological test for ovine epididymitis (*Brucella ovis*), carried out, with negative results, on a sample taken during the 30 day period prior to departure.]
	(2)*either*[II.2.9.	They originate from a Member State or a zone free from infection with bluetongue virus (serotypes 1-24), where no case of infection with bluetongue virus (serotypes 1-24) has been confirmed during the last 24 months in the targeted animal population and have not been vaccinated with a live vaccine against infection with bluetongue virus (serotypes 1-24) in the 60 day period before the date of movement and the requirements laid down in Article 32(1)(a), (b) or (c) or Article 32(2) of Delegated Regulation (EU) 2020/688 are fulfilled.]
	(2)*and/or*[II.2.9.	They originate from a Member State or a zone covered by the eradication programme for infection with bluetongue virus (serotypes 1-24) and the requirements laid down in Article 32(1)(a), (b) or (c) or Article 32(2) of Delegated Regulation (EU) 2020/688 are fulfilled, and they
	(2)*either*	[II.2.9.1. have been kept in a Member State or zone seasonally free from infection with bluetongue virus (serotypes 1-24) in accordance with Article 40(3) of Commission Delegated Regulation (EU) 2020/689
		(2)*either* [II.2.9.1.1. for at least 60 days prior to the date of movement]]

		(2)*and/or*	[II.2.9.1.2.	for at least 28 days prior to the date of movement and have been subjected to a serological test, with negative results, carried out on samples collected at least 28 days following the entry date of the animal into the Member State or zone seasonally free from infection with bluetongue virus (serotypes 1-24)]]
		(2)*and/or*	[II.2.9.1.3.	for at least 14 days prior to the date of movement and have been subjected to a PCR test, with negative results, carried out on samples collected at least 14 days following the entry date of the animal into the Member State or zone seasonally free from infection with bluetongue virus (serotypes 1-24);]]]
	(2)*and/or*	[II.2.9.2.	have been protected against attacks by the vectors during transportation to the place of destination and have been kept protected against attacks by vectors in a vector protected establishment	
		(2)*either*	[II.2.9.2.1.	for at least 60 days prior to the date of movement]]
		(2)*and/or*	[II.2.9.2.2.	for at least 28 days prior to the date of movement and have been subjected to a serological test, with negative results, carried out on samples collected at least 28 days following the date of the commencement of the period of protection against attacks by vectors]]
		(2)*and/or*	[II.2.9.2.3.	for at least 14 days prior to the date of movement and have been subjected to a PCR test, with negative results, carried out on samples collected at least 14 days following the date of the commencement of the period of protection against attacks by vectors;]]]
	(2)*and/or*	[II.2.9.3.	have been vaccinated against those serotypes from 1 to 24 of infection with bluetongue virus which were reported during the past 2 years in that Member State or zone and are within the immunity period guaranteed in the specifications of the vaccine and	
		(2)*either*	[II.2.9.3.1.	have been vaccinated more than 60 days before the date of movement]]
		(2)*and/or*	[II.2.9.3.2.	have been vaccinated with an inactivated vaccine and subjected to a PCR test, with negative results on samples collected at least 14 days after the onset of the immunity set in the specifications of the vaccine;]]]
	(2)*and/or*	[II.2.9.4.	have been subjected with positive results to a serological test able to detect specific antibodies against all serotypes 1-24 of infection with bluetongue virus reported during the past 2 years in that Member State or zone and	
		(2)*either*	[II.2.9.4.1.	the serological test has been carried out on samples collected at least 60 days before the date of movement]]
		(2) *and/or*	[II.2.9.4.2.	the serological test has been carried out on samples collected at least 30 days before the date of the movement and the animal has been subjected to a PCR test, with negative results, carried out on samples collected not earlier than 14 days before the date of movement;]]]

	[(2)]*and/or*[II.2.9.	They originate from a Member State or a zone neither free from infection with bluetongue virus (serotypes 1-24) nor covered by the eradication programme for infection with bluetongue virus (serotypes 1-24) and the requirements laid down in Article 32(1)(a), (b) or (c) or Article 32(2) of Delegated Regulation (EU) 2020/688 are fulfilled, and they			
	[(2)]*either*	[II.2.9.1.	have been protected against attacks by the vectors during transportation to the place of destination and have been kept protected against attacks by vectors in a vector protected establishment		
		[(2)]*either*	[II.2.9.1.1.	for at least 60 days prior to the date of movement]]	
		[(2)]*and/or*	[II.2.9.1.2.	for at least 28 days prior to the date of movement and have been subjected to a serological test, with negative results, carried out on samples collected at least 28 days following the date of the commencement of the period of protection against attacks by vectors]]	
		[(2)]*and/or*	[II.2.9.1.3.	for at least 14 days prior to the date of movement and have been subjected to a PCR test, with negative results, carried out on samples collected at least 14 days following the date of the commencement of the period of protection against attacks by vectors;]]]	
	[(2)]*and/or*	[II.2.9.2.	have been kept for the 60 day period prior to departure in an establishment situated in a Member State or in an area of at least 150 km radius centred on the establishment, where surveillance in compliance with the requirements set out in Sections 1 and 2 of Chapter 1 of Part II of Annex V to Delegated Regulation (EU) 2020/689 has been carried out during that period, and		
		[(2)]*either*	[II.2.9.2.1.	the animals have been vaccinated against those serotypes from 1 to 24 of infection with bluetongue virus which were reported during the past 2 years in an area of at least 150 km radius centred on the place where the animals were kept and are within the immunity period guaranteed in the specifications of the vaccine and	
			[(2)]*either*	[II.2.9.2.1.1.	have been vaccinated more than 60 days before the date of movement]]]
			[(2)]*and/or*	[II.2.9.2.1.2.	have been vaccinated with an inactivated vaccine and subjected to a PCR test, with negative results on samples collected at least 14 days after the onset of the immunity set in the specifications of the vaccine;]]]]
		[(2)]*and/or*	[II.2.9.2.2.	the animals have been immunised against those serotypes from 1 to 24 of infection with bluetongue virus which were reported during the past 2 years in an area of at least 150 km radius centred on the place where the animals were kept, and	
			[(2)]*either*	[II.2.9.2.2.1.	the animals have been subjected with positive results to a serological test carried out on samples collected at least 60 days before the date of movement]]]
			[(2)]*and/or*	[II.2.9.2.2.2.	the animals have been subjected with positive results to a serological test carried out on samples collected at least 30 days before the date of the movement and to a PCR test, with negative results, carried out on samples collected not earlier than 14 days before the date of movement;]]]]

	(2)*and/or*[II.2.9.	They do not fulfil the requirements laid down in points 1 to 3 of Section 1 of Chapter 2 of Part II of Annex V to Delegated Regulation (EU) 2020/689 and the competent authority of the Member State of origin authorised movement of those animals to another Member State or zone thereof	
	(2)*either*	[II.2.9.1.	with the status free from infection with bluetongue virus (serotypes 1-24) and the Member State of destination has informed the Commission and the other Member States that such movement is authorised subject to the conditions referred to in Article 43(2)(a), (b) and (c) of Delegated Regulation (EU) 2020/689, and
		(2)*either* [II.2.9.1.1.	point 5 of Section 1 of Chapter 2 of Part II of Annex V to that Delegated Regulation, and
		(2)*and/or* [II.2.9.1.2.	point 6 of Section 1 of Chapter 2 of Part II of Annex V to that Delegated Regulation, and
		(2)*and/or* [II.2.9.1.3.	point 7 of Section 1 of Chapter 2 of Part II of Annex V to that Delegated Regulation, and
		(2)*and/or* [II.2.9.1.4.	point 8 of Section 1 of Chapter 2 of Part II of Annex V to that Delegated Regulation, and the requirements laid down in Article 32(1)(a), (b) or (c) or Article 32(2) of Delegated Regulation (EU) 2020/688 and the requirements laid down in Article 33 of that Delegated Regulation are fulfilled.]]]
	(2)*and/or*	[II.2.9.2.	with an approved eradication program for infection with bluetongue virus (serotypes 1-24) and the Member State of destination has informed the Commission and the other Member States that such movement is authorised under the conditions referred to in Article 43(2)(a), (b) and (c) of Delegated Regulation (EU) 2020/689, and
		(2)*either* [II.2.9.2.1.	point 5 of Section 1 of Chapter 2 of Part II of Annex V to that Delegated Regulation, and
		(2)*and/or* [II.2.9.2.2.	point 6 of Section 1 of Chapter 2 of Part II of Annex V to that Delegated Regulation, and
		(2)*and/or* [II.2.9.2.3.	point 7 of Section 1 of Chapter 2 of Part II of Annex V to that Delegated Regulation, and
		(2)*and/or* [II.2.9.2.4.	point 8 of Section 1 of Chapter 2 of Part II of Annex V to that Delegated Regulation, and the requirements laid down in Article 32(1)(a), (b) or (c) or Article 32(2) of Delegated Regulation (EU) 2020/688 and the requirements laid down in Article 33 of that Delegated Regulation are fulfilled.]]]
	(2)*and/or*	[II.2.9.3.	neither free from infection with bluetongue virus (serotypes 1-24) nor covered by the eradication programme for infection with bluetongue virus (serotypes 1-24) and the Member State of destination has informed the Commission and the other Member States that such movement is authorised
		(2)*either* [II.2.9.3.1.	without any conditions, and
		(2)*and/or* [II.2.9.3.2.	subject to the conditions referred to in point 5 of Section 1 of Chapter 2 of Part II of Annex V to Delegated Regulation (EU) 2020/689, and

		(2)*and/or*	[II.2.9.3.3.	under the conditions referred to in point 6 of Section 1 of Chapter 2 of Part II of Annex V to Delegated Regulation (EU) 2020/689, and
		(2)*and/or*	[II.2.9.3.4.	under the conditions referred to in point 7 of Section 1 of Chapter 2 of Part II of Annex V to Delegated Regulation (EU) 2020/689, and
		(2)*and/or*	[II.2.9.3.5.	under the conditions referred to in point 8 of Section 1 of Chapter 2 of Part II of Annex V to Delegated Regulation (EU) 2020/689, and the requirements laid down in Article 32(1)(a), (b) or (c) or Article 32(2) of Delegated Regulation (EU) 2020/688 and the requirements laid down in Article 33 of that Delegated Regulation are fulfilled.]]]
	(2)*either* [II.2.10.	The animals are intended for a Member State or zone of a Member State listed in point 2.3. of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001 of the European Parliament and of the Council as having a negligible risk status for classical scrapie or for a Member State listed in point 3.2. of that Section as having an approved national scrapie control programme, and		
		(2)*either*	[come from a holding situated in a Member State or zone of a Member State listed in point 2.3. of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001 as having a negligible risk status for classical scrapie.]	
		(2)*and/or*	[come from a holding recognised as having a negligible risk of classical scrapie in accordance with point 1.2. of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001 and listed as such by the competent authority of the Member State in accordance with point 1.1. of that Section.]	
		(2)*and/or*	[come from a holding not subject to the measures laid down in points 3 and 4 of Chapter B of Annex VII to Regulation (EC) No 999/2001 and the animals are of the ovine species and are of the ARR/ARR prion protein genotype, or the animals are of the caprine species and carry at least one of the K222, D146 or S146 alleles.]	
		(2)*and/or*	[come from and are destined for an approved body, institute or centre as defined in Article 2(1)(c) of Council Directive 92/65/EEC.]	
		(2)*or*	[comply with the conditions set out in point 4.1.(d) of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001.]]	
	(2)*or* [II.2.10.	The animals are for breeding and are intended for a Member State or zone of a Member State other than those listed in point 2.3. of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001 as having a negligible risk status for classical scrapie or other than those listed in point 3.2. of that Section as having an approved national scrapie control programme, and		
		(2)*either*	[come from a holding situated in a Member State or zone of a Member State listed in point 2.3. of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001 as having a negligible risk status for classical scrapie.]	

(2)*and/or* [come from a holding recognised as having a negligible risk of classical scrapie in accordance with point 1.2. of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001 and listed as such by the competent authority of the Member State in accordance with point 1.1. of that Section.]

(2)*and/or* [come from a holding recognised as having a controlled risk of classical scrapie in accordance with point 1.3. of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001 and listed as such by the competent authority of the Member State in accordance with point 1.1. of that Section.]

(2)*and/or* [come from a holding not subject to the measures laid down in points 3 and 4 of Chapter B of Annex VII to Regulation (EC) No 999/2001 and the animals are of the ovine species and are of the ARR/ARR prion protein genotype, or the animals are of the caprine species and carry at least one of the K222, D146 or S146 alleles.]

(2)*and/or* [come from and are destined for an approved body, institute or centre as defined in Article 2(1)(c) of Directive 92/65/EEC.]

(2)*or* [comply with the conditions set out in point 4.1.(d) of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001.]]

(2)*or* [II.2.10. The animals are not for breeding and are intended for a Member State or zone of a Member State other than those listed in point 2.3. of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001 as having a negligible risk status for classical scrapie or other than those listed in point 3.2. of that Section as having an approved national scrapie control programme.]

II.3. To the best of my knowledge and as declared by the operator, the animals come from establishments where there were no abnormal mortalities with an undetermined cause.

(2)[II.4. According to official information and as declared by the operator, they are semen donor animals, and

II.4.1. they come from a semen collection centre and will be transported directly to another semen collection centre in accordance with Article 19 of Commission Delegated Regulation (EU) 2020/686; and

(2)*either* [II.4.2. they were continuously present since the date of their admission at the semen collection centre and were subjected, with negative results, to all compulsory routine tests referred to in point 2 of Chapter I of Part 3 of Annex II to Delegated Regulation (EU) 2020/686 in the period of the preceding 12 months prior to date of that movement; and]

(2)*or* [II.4.2. they were subjected, with negative results, to all tests referred to in point 1(c) and (d) of Chapter I of Part 3 of Annex II to Delegated Regulation (EU) 2020/686, required before admission to a semen collection centre carried out during the period immediately preceding quarantine and during the quarantine period; and]

II.4.3. the prior consent of the centre veterinarian of the semen collection centre of destination has been obtained by the operator; and

II.4.4. the means of transport used have been cleansed and disinfected before use.]

II.5. Arrangements are made to transport the consignment in accordance with Article 4 of Delegated Regulation (EU) 2020/688.

II.6. This certificate is valid for 10 days from the date of issuing. In the case of transport by waterway/sea of animals, the period of validity of the certificate may be extended by the duration of the journey by waterway/sea.

(2)(3)[II.7. Since leaving their establishments of origin and before arriving to this establishment approved for assembly operations, none of the animals of the consignment has undergone more than two assembly operations, and

(2)*either* [they come from their establishments of origin.]]

(2)*or* [at least one of the animals of the consignment has undergone one assembly operation on an approved establishment.]]

(2)*or* [at least one of the animals of the consignment has undergone two assembly operations on approved establishments.]]

Animal welfare attestation

At the time of inspection, the animals covered by this health certificate were fit to be transported in accordance with the provisions of Council Regulation (EC) No 1/2005 on the intended journey due to start on……*(insert date)* (4)(5).

Notes:

In accordance with the Agreement on the withdrawal of the United Kingdom of Great Britain and Northern Ireland from the European Union and the European Atomic Energy Community, and in particular Article 5(4) of the Protocol on Ireland / Northern Ireland in conjunction with Annex 2 to that Protocol, references to European Union in this certificate include the United Kingdom in respect of Northern Ireland.

This animal health certificate shall be completed according to the notes for the completion of certificates provided for in Chapter 2 of Annex I to Commission Implementing Regulation (EU) 2020/2235.

Part I:

Box reference I.11: "*Place of dispatch":* Indicate an establishment of the origin of the animals in the consignment or an establishment approved for assembly operations in accordance with Articles 97 and 99 of Regulation (EU) 2016/429 of the European Parliament and of the Council.

Box reference I.12: "*Place of destination":* Indicate an establishment of the final destination of the consignment or an establishment approved for assembly operations in accordance with Articles 97 and 99 of Regulation (EU) 2016/429.

Box reference I.17: "*Accompanying documents"*: In case the animals are dispatched from an establishment approved for assembly operations in the Member State of origin, the reference number(s) of the official document(s), based on which the animal health certificate for this consignment is issued in this establishment approved for assembly operations, may be indicated.

In case the animals are dispatched from an establishment approved for assembly operations in the Member State of passage, the reference number(s) of the certificate(s), based on which the animal health certificate for this consignment is issued in this establishment approved for assembly operations, must be indicated.

Box reference I.30: "*Identification number"*: Indicate identification codes of the animals in the consignment identified in accordance with Article 45(2) or (4) or Article 46(1) of Delegated Regulation (EU) 2019/2035.

Part II:

(1) There can be one or more animals in the consignment.

(2) Delete if not applicable.

(3) Applicable in case the consignment is dispatched from the establishment approved for assembly operations.

(4) This statement does not exempt transporters from their obligation in accordance with Union rules in force in particular regarding the fitness to be transported.

(5) To be completed in case of consignment grouped in an establishment approved for assembly operations located in the Member State of transit.

Official veterinarian

Name (in capital letters) Qualification and title

Local Control Unit name Local Control Unit code

Date

Stamp Signature