

Part 2

**SEMEN OF DOMESTIC ANIMALS OF THE BOVINE SPECIES
COLLECTED, PROCESSED AND STORED BEFORE 31 DECEMBER 2004 FOR
IMPORT FROM 1 JANUARY 2005 IN ACCORDANCE WITH ARTICLE 2(2) OF
COUNCIL DIRECTIVE 2003/43/EC**

Parti 2

**SEMEN TA' ANNIMALI DOMESTIČI TA' L-ISPEČI BOVINA
MIĠBUR, IPPROĊESSAT U MAĦŽUN QABEL IL-31 TA' DIĊEMBRU 2004 GHALL-
IMPORTAZZJONI MILL-1 TA' JANNAR 2005 SKOND L-ARTIKOLU 2(2) TAD-
DIRETTIVA TAL-KUNSILL 2003/43/KE**

The following model certificate is applicable from 1 January 2005 to imports of stocks of semen collected, processed and stored before 31 December 2004 in accordance with the conditions previously laid down in Council Directive 88/407/EEC and imported after that date in accordance with Article 2(2) of Directive 2003/43/EC.

Iċ-ċertifikat mudell li ġej huwa applikabbli mill-1 ta' Jannar 2005 għall-importazzjoni ta' provvisti ta' semen miġbur, ipproċessat u maħżun qabel il-31 ta' Diċembru 2004 skond il-kondizzjonijiet stabbiliti qabel fid-Direttiva tal-Kunsill 88/407/KEE u importat wara dik id-data skond l-Artikolu 2(2) tad-Direttiva 2003/43/KE.

COUNTRY

Veterinary certificate to EU

Part I : Details of dispatched consignment	1.1. Consignor <input type="checkbox"/> Name Address Postal code		1.2.		1.2.a. Local reference number:	
	1.5. Consignee Name Address Postal code		1.3. Central Competent Authority			
	1.7. Country of origin		ISO code	1.8. Region of origin		Code
	1.9. Country of destination		ISO code	1.10. Region of destination		Code
	1.11. Place of origin Semen centre <input type="checkbox"/> Name Address Approval number Name Address Approval number Name Address Approval number		1.12. Place of destination Holding <input type="checkbox"/> Semen centre <input type="checkbox"/> Approved body <input type="checkbox"/> Name Address Approval number Postal code			
	1.13.		1.14. Estimated date and time of arrival			
	1.15. Means of transport Aeroplane <input type="checkbox"/> Ship <input type="checkbox"/> Railway wagon <input type="checkbox"/> Road vehicle <input type="checkbox"/> Other <input type="checkbox"/> Identification: Documentary references:		1.16.			
	1.18. Description of commodity		1.19. Commodity code (HS code)		1.20. Quantity	
	1.21.		1.22. Number of packages			
	1.23. Identification of container/Seal number		1.24.			
	1.25. Commodity certified for Artificial reproduction <input type="checkbox"/>					
	1.26. For transit to 3rd Country vis-à-vis EU 3rd country		ISO code	1.27. For import or admission into EU Definitive import		<input type="checkbox"/>
	1.28. Identification of the animals/products Species (Scientific name) Identification mark Quantity of doses Approval number of the centre of origin					

PAJJIZ

Ic-certifikat veterinarju ghall-UE

Parti I : Dettalji tal-kunsinjja mibghuta	I.1. Kunsinatur <input type="checkbox"/> Isem		I.2.		I.2.a. Numru lokali ta' riferenza			
	Indirizz Kodici Postali		I.3. Awtorita Centrali kompetenti					
			I.4. Awtorita lokali kompetenti					
	I.5. Destinatarju Isem Indirizz Kodici Postali		I.6.					
	I.7. Pajjiz ta' origini	Kodici ISO	I.8. Regjun ta' origini	Kodici	I.9. Pajjiz ta' destinazzjoni	Kodici ISO	I.10. Regjun ta' destinazzjoni	Kodici
	I.11. Pajjiz ta' origini/Pajjiz ta' hsd Centru ta' semen <input type="checkbox"/>				I.12. Post ta' destinazzjoni			
	Isem		Numru approvat		Fond <input type="checkbox"/>		Centru ta' semen <input type="checkbox"/>	Korp approvat <input type="checkbox"/>
	Indirizz				Isem		Numru approvat	
	Isem		Numru approvat		Indirizz			
	Indirizz				Kodici Postali			
	Isem		Numru approvat		I.14. Estimazzjoni tad-data u l-hin tal-wasla			
	Indirizz							
	I.13.							
	I.15. Mezz ta' trasport Ajruplan <input type="checkbox"/> Vaxxel <input type="checkbox"/> Vagun <input type="checkbox"/> Vejkolu tat-triq <input type="checkbox"/> Ohrajn <input type="checkbox"/>				I.16.			
Identifikazzjoni Referenzi Dokumentarji:				I.17.				
I.18. Deskrizzjoni tal-prodott					I.19. Kodici tal-Komodita (Kodici CN)			
					I.20. Numru/Kwantita			
I.21.					I.22. Numru ta' pakketti			
I.23. Identifikazzjoni tal-kontenitur/ Numru ta' sigill					I.24.			
I.25. Prodotti ccertifikati Riproduzzjoni artifizjali <input type="checkbox"/>								
I.26. Ghat-tranzitu fl-UE lejn Pajjiz Terz Pajjiz terz				I.27. Ghall-importazzjoni jew l-ammissjoni fl-UE Importazzjoni definitiva				
Kodici ISO								
I.28. Identifikazzjoni tal-prodotti Speci (Isem xjentifiku) Marka ta' identifikazzjoni Numru ta' dozi Numru ta' approvazzjoni tac-centru ta' origini								

COUNTRY

Domestic bovine semen collected, processed
and stored before 31 December 2004

PAJJIŻ

Semen ta' annimali bovini domestiċi miġbur
ipproċessat u maħżun qabel il-31ta'Diċembru 2004

Part II: Certification/ Partii II: Ċertifikazzjoni	<p>II. Health information <input type="checkbox"/></p> <p><i>Taghrif fuq is-sahha</i> <input type="checkbox"/></p>	<p>II.a. Certificate reference number</p> <p><i>Numru ta' referenza taċ-Ċertifikat</i></p>	<p>II.b. Local reference number</p> <p><i>Numru ta' referenza lokali</i></p>
	<p>I, the undersigned, official veterinarian, hereby certify that:</p> <p>1.1. (Name of exporting country) ⁽⁴⁾</p> <p>was free from rinderpest and foot-and-mouth disease during the 12 months immediately prior to collection of the semen for export and up until its date of dispatch and no vaccination against these diseases took place during that period;</p> <p><i>Jiena, il-veterinarju uffiċjali hawn taht iffirmat, niċċertifika li:</i></p> <p>..... (Isem il-pajjiż esportatur) ⁽³⁾</p> <p><i>kien hieles mir-rinderpest u l-marda ta' l-ilsien u d-dwiefer matul it-12-11 xahar immedjatament qabel il-ġbir tas-semen għall-esportazzjoni u sad-data tal-bgħit tiegħu u ma sar l-ebda tilqim kontra dan il-mard matul dak il-perjodu;</i></p> <p>1.2. The semen described above was collected before 31 December 2004 at a semen collection centre which: <i>Is-semen deskritt hawn fuq ingabar qabel il-31 ta' Diċembru 2004 f'ċentru ta' ġbir tas-semen li:</i></p> <p>1.2.1. meets the conditions laid down in Chapter I of Annex A to Directive 88/407/EEC; <i>jissodisfa l-kondizzjonijiet stabbiliti fil-Kapitolu I ta' l-Anness A għad-Direttiva 88/407/KEE;</i></p> <p>1.2.2. is operated and supervised in accordance with the conditions laid down in Chapter II of Annex A to Directive 88/407/EEC; <i>huwa mħaddem u ssorveljat skond il-kondizzjonijiet stabbiliti fil-Kapitolu II ta' l-Anness A għad-Direttiva tal-Kunsill 88/407/KEE;</i></p> <p>1.3. The centre at which the semen to be exported was collected was free from rabies, tuberculosis, brucellosis, anthrax and contagious bovine pleuropneumonia during the 30 days prior to the date of collection of the semen to be exported and the 30 days after collection (in the case of fresh semen, until the date of dispatch); <i>Iċ-ċentru fejn ingabar is-semen għall-esportazzjoni, kien hieles mill-idrofobija/rabja, tuberkulozi, brucellozi, antraċe u plewropnewmonja bovina kontaġjuża matul it-30 jum qabel id-data tal-ġbir tas-semen għall-esportazzjoni u t-30 jum wara l-ġbir (fil-każ ta' semen frisk, sad-data tal-bgħit);</i></p> <p>1.4. At the time the semen described above was collected, all bovine animals at the semen collection centre: <i>Fil-ħin tal-ġbir tas-semen deskritt hawn fuq, l-annimali bovini kollha fiċ-ċentru ta' ġbir tas-semen:</i></p>		

	<p>1.4.1. came from herds and/or were born to dams which satisfy the conditions in paragraph 1(b) and (c) of Chapter I of Annex B to Directive 88/407/EEC;</p> <p><i>għew minn merħliet u/jew twieldu minn ommijiet li jissodisfaw il-kondizzjonijiet fil-paragrafu 1(b) u (c) tal-Kapitolu I ta' l-Anness B għad-Direttiva 88/407/KEE;</i></p> <p>1.4.2. had tested negative, within the 30 days preceding the quarantine isolation period, to:</p> <p>the tests referred to in points 1(d)(i), (ii) and (iii) of Chapter I of Annex B to Directive 88/407/EEC, and</p> <ul style="list-style-type: none"> – a serum neutralisation test or an ELISA test for infectious bovine rhinotracheitis/infectious pustular vulvo-vaginitis, and – a virus isolation test (fluorescent antibody test or immunoperoxidase test) for bovine viral diarrhoea, deferred until the animal reached the age of six months in the case of younger animals; <p><i>irriżultaw negattivi, matul it-30 jum qabel il-perjodu ta' iżolazzjoni tal-kwarantina, għat-:</i></p> <ul style="list-style-type: none"> – <i>testijiet imsemmija f'punti 1(d)(i), (ii) u (iii) tal-Kapitolu I ta' l-Anness B għad-Direttiva 88/407/KEE, u</i> – <i>test tan-newtralizzazzjoni tas-serum jew test ELISA għar-rinotrakejite bovina infettiva/pustular vulvo-vaginitis infettiva, u</i> – <i>test għall-iżolazzjoni tal-vajrus (test florixxenti ta' l-antikorpi jew test immunoperoxidase) għal dijarea virali fil-bovini, differit sakemm l-annimal jilhaq is-sitt xhur fil-każ ta' annimali iżgħar fiż-żmien;</i> <p>1.4.3. had undergone the 30-day quarantine isolation period and had tested negative to the following health tests:</p> <ul style="list-style-type: none"> – a serological test for brucellosis carried out in accordance with the procedure described in Annex C to Directive 64/432/EEC, – either an immunofluorescent antibody test or a culture test for campylobacter fetus infection on a sample of preputial material or artificial vagina washings or, in the case of a female animal, a vaginal mucus agglutination test⁽¹⁾, – a microscopic examination and culture test for trichomonas foetus on a sample of preputial material or artificial vagina washings or, in the case of a female animal, a vaginal mucus agglutination test⁽¹⁾; <p><i>kienu għamlu t-30 jum f'izolazzjoni ta' kwarantina u kienu rriżultaw negattivi għat-testijiet ta' saħħa li ġejjin:</i></p> <ul style="list-style-type: none"> – <i>test serologiku għall-brucellożi mwettaq skond il-proċedura deskritta fl-Anness Ċ għad-Direttiva 64/432/KEE,</i> – <i>jew test ta' immunoflorixxenza ta' l-antikorpi jew test ta' kultura għall-infezzjoni campylobacter tal-fetu fuq kampjun ta' materjal perpużjali jew hasliet ta' vagina artifiċjali jew, fil-każ ta' annimali femminili, test ta' agglutinazzjoni tal-mukus vaġinali⁽¹⁾,</i> – <i>eżami mikroskopiku u test ta' kultura għat-trichomonas tal-fetu fuq kampjun ta' materjal prupuzjali jew hasliet tal-vaġina jew, fil-każ ta' annimali femminili, test ta' agglutinazzjoni tal-mukus vaġinali⁽¹⁾;</i> <p>1.4.4. had tested negative, at least once a year, to the routine tests referred to in points 1(a), (b) and (c) of Chapter II of Annex B to Directive 88/407/EEC;</p> <p><i>kienu rriżultaw negattivi, ta' l-inqas darba f'sena, għat-testijiet ta' rutina msemmija fil-punti 1(a), (b) u (c) tal-Kapitolu II ta' l-Anness B għad-Direttiva 88/407/KEE;</i></p> <p>1.5. At the time the semen described above was collected,</p> <p><i>Fiż-żmien li fih ingabar is-semen deskritt hawn fuq,</i></p> <p>1.5.1. all female bovine animals in the centre had tested negative at least once a year to a vaginal mucus agglutination test for campylobacter fetus infection, and</p> <p><i>l-annimali femminili kollha fiċ-ċentru irriżultaw negattivi ta' l-inqas darba f'sena għal test ta' agglutinazzjoni tal-mukus vaġinali għall-infezzjoni campylobacter tal-fetu, u</i></p>
--	---

1.5.2. all bulls used for semen production had tested negative either to an immunofluorescent antibody test or to a culture test for campylobacter fetus infection on a sample of preputial material or artificial vagina washings carried out in the 12 months prior to collection;

il-barrin kollha użati għall-produzzjoni tas-semen irriżultaw negattivi jew għal test ta' immunofloressenza ta' l-antikorpi jew għal test ta' kultura għall-infezzjoni campylobacter tal-fetu fuq kampjun ta' materjal prepuzjali jew hasliet ta' vagina artifiċjali mwettqa fit-12-il xahar qabel il-ġbir;

1.6. The semen to be exported was obtained from donor bulls which:

Is-semen għall-esportazzjoni inkiseb minn barrin donaturi li:

1.6.1. satisfy the conditions laid down in Annex C to Directive 88/407/EEC;

jissodisfaw il-kondizzjonijiet stabbiliti fl-Anness Ċ għad-Direttiva 88/407/KEE;

1.6.2. either were resident in the exporting country during the six months immediately prior to collection of the semen for export⁽¹⁾;

or

had been imported from.....⁽⁴⁾, after spending less than six months in the exporting country, and at the time of import, satisfied the health conditions applying to donors the semen of which is intended for export to the Community⁽¹⁾;

jew kienu residenti fil-pajjiż esportatur matul is-sitt xhur immedjatament qabel il-ġbir tas-semen għall-esportazzjoni⁽¹⁾;

jew

ġew importati minn⁽³⁾, wara li qattgħu inqas minn sitt xhur fil-pajjiż esportatur, u fil-hin ta' l-importazzjoni, issodisfaw il-kondizzjonijiet ta' saħħa applikabbli għad-donaturi li s-semen tagħhom huwa maħsub għall-esportazzjoni lejn il-Komunità⁽¹⁾;

1.6.3. stand in a semen collection centre at which:

- (i) all bovine animals tested negative at least once a year to a serum neutralisation test or an ELISA test for infectious bovine rhinotracheitis/infectious pustular vulvo-vaginitis⁽¹⁾, or
- (ii) bovine animals not vaccinated against infectious bovine rhinotracheitis tested negative at least once a year to a serum neutralisation test or an ELISA test for infectious bovine rhinotracheitis/infectious pustular vulvo-vaginitis and at which testing for infectious bovine rhinotracheitis was not carried out on bulls which had received their first vaccination against infectious bovine rhinotracheitis at the insemination centre after they had tested negative to a serum neutralisation test or an ELISA test for infectious bovine rhinotracheitis/infectious pustular vulvo-vaginitis and which had been regularly re-vaccinated at intervals of not more than six months since the first vaccination⁽¹⁾;

jinsabu f'ċentru ta' ġbir tas-semen fejn:

- i) *l-annimali bovini kollha rriżultaw negattivi ta' l-inqas darba fis-sena għal test ta' newtralizzazzjoni tas-serum jew test ELISA għar-rinotrakejite bovina infettiva/pustular vulvo-vaginitis infettiva⁽¹⁾, jew*
- ii) *l-annimali bovini li mhumiex imlaqqma kontra r-rinotrakejite bovina infettiva rriżultaw negattivi ta' l-inqas darba fis-sena għal test ta' newtralizzazzjoni tas-serum jew test ELISA għar-rinotrakejite bovina infettiva/pustular vulvo-vaginitis infettiva u li fihom ma sarux testijiet għar-rinotrakejite bovina infettiva fuq barrin li kienu hađu l-ewwel tilqima kontra r-rinotrakejite bovina infettiva fiċ-ċentru ta' inseminazzjoni wara li kienu rriżultaw negattivi għal test tan-newtralizzazzjoni tas-serum jew test ELISA għar-rinotrakejite bovina infettiva/pustular vulvo-vaginitis infettiva u li kienu mlaqqma regolarmnt f'intervalli ta' mhux iktar minn sitt xhur mill-ewwel tilqima⁽¹⁾;*

1.6.4. fulfil the import conditions for bovine semen laid down in the Bluetongue Chapter of the Terrestrial Animal Health Code of the OIE, depending on the status of the country or zone of residence; ****

*jissodisfaw il-kondizzjonijiet ta' importazzjoni ta' semen bovin stabbiliti fil-Kapitolu tal-Bluetongue tal-Kodiċi tas-Saħħa ta' l-Annimali Terrestri ta' l-OIE, skond l-istatus tal-pajjiż jew żona ta' fejn jinżammu; *****

1.6.5. were resident in the country of export in which the following serotypes of epizootic haemorrhagic disease (EHD) exist:; and tested negative on two occasions not more than 12 months apart to an agar-gel immuno-diffusion test⁽⁵⁾ and a virus neutralisation test for all above-listed serotypes of EHD, carried out in an approved laboratory on samples of blood taken prior to and not less than 21 days following collection of the semen; ***

*kienu residenti fil-pajjiż esportatur fejn jeżistu s-serotipi tal-marda epizootika emorraġika (EHD) li ġejjin:; u rriżultaw negattivi f'żewġ okkażjonijiet mhux aktar minn 12-il xahar bogħod minn xulxin għal test ta' immuno-diffużjoni bl-agar ġel⁽⁴⁾ u test ta' newtralizzazzjoni tal-vajrus tas-serotipi kollha ta' l-EHD elenkati hawn fuq, imwettqa f'laboratorju approvat fuq kampjuni tad-demem mehuda qabel il-ġbir tas-semen u mhux inqas minn 21 jum wara;****

1.6.6. were resident in the country of export in which the following serotypes of epizootic haemorrhagic disease (EHD) exist:; and tested negative, prior to entry and at six-monthly intervals, to an agar-gel immuno-diffusion test⁽⁵⁾ and a virus neutralisation test for all above-listed serotypes of EHD, carried out in an approved laboratory; **

*kienu residenti fil-pajjiż esportatur fejn jeżistu s-serotipi tal-marda epizootika emorraġika (EHD) li ġejjin:; u rriżultaw negattivi, qabel id-dhul u f'perjodi ta' sitt xhur, għal test ta' l-immuno-diffużjoni bl-agar ġel⁽⁴⁾ u għal test ta' newtralizzazzjoni tal-vajrus tas-serotipi kollha ta' l-EHD elenkati hawn fuq, imwettqa f'laboratorju approvat; ***

1.6.7. tested negative on two occasions not more than 12 months apart to a serum neutralisation test for Akabane virus, carried out in an approved laboratory on a blood sample taken prior to and not less than 21 days following collection of the semen; *

*irriżultaw negattivi f'żewġ okkażjonijiet mhux aktar minn 12-il xahar bogħod minn xulxin għal test ta' newtralizzazzjoni tas-serum għall-vajrus Akabane, imwettqa f'laboratorju approvat fuq kampjuni tad-demem mehuda qabel il-ġbir tas-semen u mhux inqas minn 21 jum wara ;**

1.7. The semen to be exported was collected after the date on which the centre was approved by the competent national authorities of the exporting country;

Is-semen għall-esportazzjoni inġabar wara d-data ta' l-approvazzjoni tač-ċentru mill-awtoritajiet nazzjonali kompetenti tal-pajjiż esportatur;

1.8. The semen to be exported was processed, stored and transported under conditions which satisfy the terms of Directive 88/407/EEC prior to its amendment by Directive 2003/43/EC.

Is-semen għall-esportazzjoni ġie pproċessat, maħżun u ttrasportat taht kondizzjonijiet li jissodisfaw it-termini tad-Direttiva 88/407/KEE qabel l-emenda tagħha bid-Direttiva 2003/43/KE.

Notes

Note for importer: this certificate is for veterinary purposes only and must accompany the consignment until it reaches the border inspection post.

- (1) Delete as necessary.
- (2) [Box reference no. I.28. in Part I]:
Identification mark: corresponding to the identification of the donor animals and the date of collection, that must be prior to 31 December 2004.
Approval number of the centre of origin: to be filled in if different from box reference no.I.11.
- (3) Countries listed in Annex I to Decision 2004/639/EC.
- (4) Standards for EHD virus diagnostic tests are described in the Bluetongue Chapter of the Manual of Diagnostic Tests and Vaccines for Terrestrial Animals.
**** To be used only by Australia, Canada and the USA.
*** To be used only by Australia and the USA.
** To be used only by Canada.
* To be used only by Australia.

Noti

Nota għall-importatur : dan iċ-ċertifikat huwa għal skopijiet veterinarji biss u għandu jakkumpanja l-kunsinna sakemm tasal fil-post ta' spezzjoni.

- (1) *Hassar kif mehtieg.*
- (2) *[Kaxxa bin-nru ta' referenza I.28 f'Parti I]:*
Marka ta' identifikazzjoni: li tikkorrispondi ma' l-identifikazzjoni ta' l-animali donaturi u d-data tal-gbir, li għandha tkun qabel il-31 ta' Diċembru 2004.
Numru ta' approvazzjoni taċ-ċentru ta' oriġini: biex jimtela jekk differenti mill-kaxxa bin-nru ta' referenza I.11.
- (3) *Pajjiżi elenkati fl-Anness I għad-Deċiżjoni 2004/639/KE.*
- (4) *Standards għat-testijiet dijanjostiċi tal-vajrus EHD huma deskritti fil-Kapitolu tal-Bluetongue tal-Manwal tat-Testijiet Dijanjostiċi u l-Vaċċini għall-Animali Terrestri.*
**** *Biex jintuża fl-Awstralja, il-Kanada u l-Istati Uniti biss.*
*** *Biex jintuża fl-Awstralja u l-Istati Uniti biss.*
** *Biex jintuża fil-Kanada biss.*
* *Biex jintuża fl-Awstralja biss.*

NB: This certificate must:

- (a) be drawn up in at least one official language of the Member State of destination and of the Member State where the embryos will enter Community territory;
- (b) be made out to a single consignee;
- (c) accompany the embryos in the original.

NB: Dan iċ-ċertifikat għandu:

- (a) *jithejja f'mill-inqas waħda mil-lingwi uffiċjali ta' l-Istat Membru tad-destinazzjoni u ta' l-Istat Membru li minnu s-semen jidhol fit-territorju tal-Komunità;*
- (b) *jimtela għal destinatarju wiehed;*
- (c) *jakkumpanja s-semen bil-kopja oriġinali.*

	<p>Official veterinarian</p> <p>Name (in Capital): Date: Stamp</p>	<p>Qualification and title Signature:</p>
	<p><i>Veterinarju ufficiali</i></p> <p><i>Isem (Ittri kbar):</i> <i>Data:</i> <i>Timbru</i></p>	<p><i>Kwalifika u Titlu</i> <i>Firma:</i></p>