Council Directive 92/118/EEC of 17 December 1992 laying down animal health and public health requirements governing trade in and imports into the Community of products not subject to the said requirements laid down in specific Community rules referred to in Annex A (I) to Directive 89/662/EEC and, as regards pathogens, to Directive 90/425/EEC

CHAPTER I

General provisions

	General provisions
Article 1 Article 2 Article 3	This Directive lays down the animal health and public health (1) For the purposes of this Directive: Member States shall ensure that: trade in and imports of
	CHAPTER II
	Provisions applicable to trade
Article 4 Article 5	Member States shall take the necessary measures to ensure that, Member States shall ensure that every necessary measure is taken
Article 6 Article 7	Member States shall ensure that trade in pathogenic agents is (1) The rules on checks established by Directive 89/662/EEC and,
Article 8	In Chapter 1 (1) of Annex A to Directive 92/46/EEC
	CHAPTER III
	Provisions applicable to imports into the Community
Article 9	The requirements applicable to imports of products covered by this
Article 10 Article 11 Article 12	(1) For the purposes of uniform application of Article 9, The procedure provided for in Article 18 shall be used (1) The principles and rules laid down in Directives 90/675/EEC
Article 13	(1) Member States may, by issuing an appropriate licence, permit
	CHAPTER IV
	Common final provisions
Article 14 Article 15 Article 16 Article 17	 Article 3 (d) of Directive 72/461/EEC shall be deleted The Council, acting by a qualified majority on a proposal Member States shall be authorized to make the entry Annexes A and B to Directives 89/662/EEC and 90/425/EEC

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	Article 18 Article 19	Where reference is made to the procedure provided for in Under the procedure provided for in Article 18, transitional
	A 4: 1 20	measures
	Article 20 Article 21	(1) Member States shall bring into force the laws, regulations This Directive is addressed to the Member States.
		ANNEX I
		SPECIFIC ANIMAL HEALTH REQUIREMENTS
		CHAPTER I
	Milk,	milk products and colostrum not intended for human consumption
		CHAPTER 2
		Animal casings intended for human consumption
A.	Trade	
B.	Imports	from third countries
		CHAPTER 3
		es and skins of ungulates not covered by Directive 64/433/EEC or 462/EEC and which have not undergone certain tanning processes
I.		462/EEC and which have not undergone certain tanning processes
I. II.	72/4	462/EEC and which have not undergone certain tanning processes
	72/4	462/EEC and which have not undergone certain tanning processes
II.	72/4	462/EEC and which have not undergone certain tanning processes
II.	72/4 Intra-Co Importa	462/EEC and which have not undergone certain tanning processes ommunity trade tions
II.	72/4 Intra-Co Importa	A62/EEC and which have not undergone certain tanning processes ommunity trade tions CHAPTER 4 ontaining low-risk materials within the meaning of Directive 90/667/EEC
II. III.	Intra-Co Importa Pet food co	A62/EEC and which have not undergone certain tanning processes ommunity trade tions CHAPTER 4 ontaining low-risk materials within the meaning of Directive 90/667/EEC
II. III.	Intra-Co Importa Pet food co	A62/EEC and which have not undergone certain tanning processes community trade tions CHAPTER 4 containing low-risk materials within the meaning of Directive 90/667/EEC

CHAPTER 5

Bones and bone products (excluding bone meal), horns and horn products (excluding horn meal) and hooves and hoof products (excluding hoof meal)intended for human consumption

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CHAPTER 6

	Processed animal proteinintended for human consumption	
I.	Without prejudice to any restrictions imposed as regards BSE or	
II.	Member States may carry out random sampling of bulk consignments	
III.	Member States must keep records of the results of sampling	
IV.	In accordance with Article 3 (3) of Directive 89/662/EEC, transhipment	
V.	Where a consignment proves to be positive for salmonella, it	
	CHAPTER 7	
	Blood and blood products of ungulates and poultry	
I. II.	Fresh blood and blood products intended for human consumption A. Trade 1. Trade in fresh blood of ungulates and poultry intended for 2. Trade in blood products intended for human consumption is subject B. Imports 1. Imports of fresh blood of domestic ungulates intended for human 2. Imports of blood products for human consumption, including those referred Fresh blood and blood products not intended for human consumption A. Definitions	
	B. Trade C. Imports 1	
III.	General provisions	
	CHAPTER 8	
	Serum from equidae	
1.		
2.		

CHAPTER 9

Lard and rendered fats intended for human consumption

- 1. Member States shall authorize the importation into the Community of...
- 2. Where there has been an outbreak of a serious transmissible...

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CHAPTER 10

Raw material for the manufacture of animal feedingstuffs and pharmaceutical or technical products

1.	
2.	
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	CHAPTER 11
	Rabbit meat and farmed game meat intended for human consumption
	CHAPTER 12
	Apiculture products
1.	
2.	
	CHAPTER 13
	Game trophies
A.	
B.	
	CHAPTER 14
	Manure I. Unprocessed manure A. Trade in unprocessed manure 1
	2

	3 B. Imports of unprocessed manure II. Processed manure and processed manure products A B III. Guano
	CHAPTER 15
	Unprocessed wool, hair, bristles, feathers and parts of feathers
1.	
2.	
3.	
4.	
	ANNEX II
	SPECIFIC PUBLIC HEALTH CONDITIONS
	CHAPTER 1
	Imports from third countries of meat products obtained from poultrymeat, farmed game meat, wild game meat and rabbit meat
	CHAPTER 2
	CHAPTER 3
I.	Specific public health conditions applicable to trade in and imports
II.	Specific public health conditions applicable to trade in and imports
	CHAPTER 4
	Section A
	SPECIFIC HEALTH CONDITIONS FOR THE GELATINE INTENDED FOR HUMAN CONSUMPTION
I.	Conditions for establishments producing gelatine
II.	Requirements for raw materials to be used for the production 1. For the production of gelatine intended for human consumption, the 2

	4.	
	5.	
	6.	
	7.	<u></u>
	8. 9.	The collection centres and tanneries which intend to supply raw Imports into the Community of raw material destined to the
III.	Transport and storage of raw materials	
	1.	
	2.	
	3.	
IV.	Condi	tions to be complied with for the manufacture of gelatine
	1.	Gelatine must be produced by a process which ensures that:
	2.	
	3. 4.	
	4.	
V.	Requi	rements for finished products
	1.	Microbiological criteria
	2.	Residues
VI.	Packa	ging, storage and transport
	1.	Gelatine intended for human consumption must be wrapped, packaged, stored
	2.	Wrappings and packages containing gelatine must:
	3.	
VII.	Importation of gelatine from third countries	
	A.	
	В.	
VIII.	Mode	l of commercial document for raw material destined to the
		Section B
		SPECIFIC HEALTH CONDITIONS FOR THE COLLAGEN INTENDED FOR HUMAN CONSUMPTION
I.	Gener	al
	1.	
	2.	
	3.	
II.	Establ	ishments producing collagen
III.	Raw n	naterials and establishments supplying them
	1.	The following raw materials may be used for the production
	2.	<u> </u>
	3.	The raw materials shall meet the following requirements:
	4.	
IV.	Transp	port and storage of the raw material
	1.	

EEC...

	2
V.	Manufacture of collagen 1
VI.	Finished products
VII.	Packaging, storage and transport 1
VIII.	Import from third countries of collagen and raw materials intended 1. Member States shall authorise import into the Community of collagen 2. Member States shall authorise import into the Community of the 3
	ANNEX III
I	CONSOLIDATED VERSION OF ANNEXES A AND B TO DIRECTIVE 89/662/EEC
II	CONSOLIDATED VERSION OF ANNEXES A AND B TO DIRECTIVE 90/425/

- (1) OJ No C 327, 30. 12. 1989, p. 29; and OJ No C 84, 2.4. 1990, p 102.
- (2) OJ No C 113, 7. 5. 1990, p. 205; and OJ No C 149, 18. 6. 1990, p. 259.
- (3) OJ No C 124, 21. 5. 1990, p. 15; and OJ No C 182, 23. 7. 1990, p. 250.
- (4) OJ No L 373, 31. 12. 1990, p. 1.