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ANNEX I

Microbiological criteria for foodstuffs

Chapter 1. Food safety criteria

Food category	Micro-organisms/ their toxins, metabolites	Sampling-plan ^a		Limits ^b		Analytical reference method ^c	Stage where the criterion applies
		n	c	m	M		
1.1.	<i>Listeria monocytogenes</i> Ready-to-eat foods intended for infants and ready-to-eat foods for special medical purposes ^d	10	0	Absence in 25 g		EN/ISO 11290-1	Products placed on the market during their shelf-life
1.2.	<i>Listeria monocytogenes</i> Ready-to-eat foods able to support the growth of <i>L. monocytogenes</i> , other than those intended for infants and for special	5	0	100 cfu/g ^e		EN/ISO 11290-2 ^f	Products placed on the market during their shelf-life
		5	0	Absence in 25 g ^g		EN/ISO 11290-1	Before the food has left the immediate control of the food business operator, who has produced it

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	medical purposes					
1.3.	Ready-to-eat foods unable to support the growth of <i>Listeria monocytogenes</i> , other than those intended for infants and for special medical purposes ^{dh}	5	0	100 cfu/g	EN/ISO 11290-2 ^f	Products placed on the market during their shelf-life
1.4.	Minced meat and meat preparations intended to be eaten raw	5	0	Absence in 25 g	EN/ISO 6579	Products placed on the market during their shelf-life
1.5.	Minced meat and meat preparations made from poultry meat intended to be eaten cooked	5	0	From 1.1.2006 Absence in 10 g From 1.1.2010 Absence in 25 g	EN/ISO 6579	Products placed on the market during their shelf-life

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1.6.	Minced meat and meat preparations made from other species than poultry intended to be eaten cooked	<i>Salmonella</i> 5	0	Absence in 10 g	EN/ISO 6579	Products placed on the market during their shelf-life
1.7.	Mechanically separated meat (MSM) ⁱ	<i>Salmonella</i> 5	0	Absence in 10 g	EN/ISO 6579	Products placed on the market during their shelf-life
1.8.	Meat products intended to be eaten raw, excluding products where the manufacturing process or the composition of the product will eliminate the salmonella risk	<i>Salmonella</i> 5	0	Absence in 25 g	EN/ISO 6579	Products placed on the market during their shelf-life
1.9.	Meat products	<i>Salmonella</i> 5	0	From 1.1.2006 Absence in 10 g From 1.1.2010	EN/ISO 6579	Products placed on the

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	made from poultry meat intended to be eaten cooked			Absence in 25 g		market during their shelf-life
1.10.	<i>Salmonella</i> 5 Gelatine and collagen	0		Absence in 25 g	EN/ISO 6579	Products placed on the market during their shelf-life
1.11.	<i>Salmonella</i> 5 Cheeses, butter and cream made from raw milk or milk that has undergone a lower heat treatment than pasteurisation ^j	0		Absence in 25 g	EN/ISO 6579	Products placed on the market during their shelf-life
1.12.	<i>Salmonella</i> 5 Milk powder and whey powder ^j	0		Absence in 25 g	EN/ISO 6579	Products placed on the market during their shelf-life
1.13.	<i>Salmonella</i> 5 Ice cream ^k , excluding products where the manufacturing	0		Absence in 25 g	EN/ISO 6579	Products placed on the market during their shelf-life

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	process or the composition of the product will eliminate the salmonella risk					
1.14.	Egg products, excluding products where the manufacturing process or the composition of the product will eliminate the salmonella risk	<i>Salmonella</i> 5	0	Absence in 25g	EN/ISO 6579	Products placed on the market during their shelf-life
1.15.	Ready- to- eat foods containing raw egg, excluding products where the manufacturing process or the composition of the product	<i>Salmonella</i> 5	0	Absence in 25 g or ml	EN/ISO 6579	Products placed on the market during their shelf-life

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	will eliminate the salmonella risk					
1.16.	Cooked crustaceans and molluscan shellfish	<i>Salmonella</i> 5	0	Absence in 25 g	EN/ISO 6579	Products placed on the market during their shelf-life
1.17.	Live bivalve molluscs and live echinoderms, tunicates and gastropods	<i>Salmonella</i> 5	0	Absence in 25g	EN/ISO 6579	Products placed on the market during their shelf-life
1.18.	Sprouted seeds (ready-to-eat) ¹	<i>Salmonella</i> 5	0	Absence in 25 g	EN/ISO 6579	Products placed on the market during their shelf-life
1.19.	Pre-cut fruit and vegetables (ready-to-eat)	<i>Salmonella</i> 5	0	Absence in 25 g	EN/ISO 6579	Products placed on the market during their shelf-life
1.20.	Unpasteurised fruit and vegetable juices (ready-to-eat)	<i>Salmonella</i> 5	0	Absence in 25 g	EN/ISO 6579	Products placed on the market during their shelf-life
1.21.	Cheeses milk powder	<i>Staphylococcus aureus</i> enterotoxins	0	Not detected in 25g	European screening method of	Products placed on the market

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	and whey powder, as referred to in the coagulase-positive staphylococci criteria in Chapter 2.2 of this Annex				the CRL for Milk ^m	during their shelf-life
1.22.	Dried infant formulae and dried dietary foods for special medical purposes intended for infants below six months of age, as referred to in the Enterobacteriaceae criterion in Chapter 2.2 of this Annex	<i>Salmonella</i> 30	0	Absence in 25 g	EN/ISO 6579	Products placed on the market during their shelf-life

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1.23.	Dried infant formulae and dried dietary foods for special medical purposes intended for infants below six months of age, as referred to in the Enterobacteriaceae criterion in Chapter 2.2 of this Annex	30	0	Absence in 10 g		ISO/DTS 22964	Products placed on the market during their shelf-life
1.24.	Live bivalve molluscs and live echinoderms, tunicates and gastropods	10	0	230 MPN/100g of flesh and intra-valvular liquid		ISO TS 16649-3	Products placed on the market during their shelf-life
1.25.	Fishery products from fish species associated with a	90	2	100 mg/kg	200 mg/kg	HPLC*	Products placed on the market during their shelf-life

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	high amount of histidine ^p						
1.26.	Histamine Fishery products which have undergone enzyme maturation treatment in brine, manufactured from fish species associated with a high amount of histidine ^p	9	2	200 mg/kg	400 mg/kg	HPLC ^r	Products placed on the market during their shelf-life

a n = number of units comprising the sample; c = number of sample units giving values over m or between m and M.

b For points 1.1-1.24 m=M.

c The most recent edition of the standard shall be used.

d Regular testing against the criterion is not useful in normal circumstances for the following ready-to-eat foods:

- those which have received heat treatment or other processing effective to eliminate *L. monocytogenes*, when recontamination is not possible after this treatment (e.g. products heat treated in their final package),
- fresh, uncut and unprocessed vegetables and fruits, excluding sprouted seeds,
- bread, biscuits and similar products,
- bottled or packed waters, soft drinks, beer, cider, wine, spirits and similar products,
- sugar, honey and confectionery, including cocoa and chocolate products,
- live bivalve molluscs.

e This criterion applies if the manufacturer is able to demonstrate, to the satisfaction of the competent authority, that the product will not exceed the limit 100 cfu/g throughout the shelf-life. The operator may fix intermediate limits during the process that should be low enough to guarantee that the limit of 100 cfu/g is not exceeded at the end of the shelf-life.

f 1 ml of inoculum is plated on a Petri dish of 140 mm diameter or on three Petri dishes of 90 mm diameter.

g This criterion applies to products before they have left the immediate control of the producing food business operator, when he is not able to demonstrate, to the satisfaction of the competent authority, that the product will not exceed the limit of 100 cfu/g throughout the shelf-life.

h Products with $pH \leq 4,4$ or $a_w \leq 0,92$, products with $pH \leq 5,0$ and $a_w \leq 0,94$, products with a shelf-life of less than five days are automatically considered to belong to this category. Other categories of products can also belong to this category, subject to scientific justification.

i This criterion applies to mechanically separated meat (MSM) produced with the techniques referred to in Chapter III, paragraph 3, in section V of Annex III to Regulation (EC) No 853/2004 of the European Parliament and of the Council of 29 April 2004 laying down specific hygiene rules for food of animal origin.

j Excluding products when the manufacturer can demonstrate to the satisfaction of the competent authorities that, due to the ripening time and a_w of the product where appropriate, there is no salmonella risk.

k Only ice creams containing milk ingredients.

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l	Preliminary testing of the batch of seeds before starting the sprouting process or the sampling to be carried out at the stage where the highest probability of finding <i>Salmonella</i> is expected.
m	Reference: Hennekinne et al., J. AOAC Internat. Vol. 86, No 2, 2003.
n	<i>E. coli</i> is used here as an indicator of faecal contamination.
o	A pooled sample comprising a minimum of 10 individual animals.
p	Particularly fish species of the families: <i>Scombridae</i> , <i>Clupeidae</i> , <i>Engraulidae</i> , <i>Coryfenidae</i> , <i>Pomatomidae</i> , <i>Scombrosidae</i> .
q	Single samples may be taken at retail level. In such a case the presumption laid down in Article 14(6) of Regulation (EC) No 178/2002, according to which the whole batch should be deemed unsafe, shall not apply.
r	References: 1. Malle P., Valle M., Bouquelet S. Assay of biogenic amines involved in fish decomposition. J. AOAC Internat. 1996, 79, 43-49. 2. Duflos G., Dervin C., Malle P., Bouquelet S. Relevance of matrix effect in determination of biogenic amines in plaice (<i>Pleuronectes platessa</i>) and whiting (<i>Merlangus merlangus</i>). J. AOAC Internat. 1999, 82, 1097-1101.

Interpretation of the test results

The limits given refer to each sample unit tested, excluding live bivalve molluscs and live echinoderms, tunicates and gastropods in relation to testing *E. coli*, where the limit refers to a pooled sample.

The test results demonstrate the microbiological quality of the batch tested⁽¹⁾.

L. monocytogenes in ready-to-eat foods intended for infants and for special medical purposes:

- satisfactory, if all the values observed indicate the absence of the bacterium,
- unsatisfactory, if the presence of the bacterium is detected in any of the sample units.

L. monocytogenes in ready-to-eat foods able to support the growth of *L. monocytogenes* before the food has left the immediate control of the producing food business operator when he is not able to demonstrate that the product will not exceed the limit of 100 cfu/g throughout the shelf-life:

- satisfactory, if all the values observed indicate the absence of the bacterium,
- unsatisfactory, if the presence of the bacterium is detected in any of the sample units.

L. monocytogenes in other ready-to-eat foods and *E. coli* in live bivalve molluscs:

- satisfactory, if all the values observed are \leq the limit,
- unsatisfactory, if any of the values are $>$ the limit.

Salmonella in different food categories:

- satisfactory, if all the values observed indicate the absence of the bacterium,
- unsatisfactory, if the presence of the bacterium is detected in any of the sample units.

Staphylococcal enterotoxins in dairy products:

- satisfactory, if in all the sample units the enterotoxins are not detected,
- unsatisfactory, if the enterotoxins are detected in any of the sample units.

Enterobacter sakazakii in dried infant formulae and dried dietary foods for special medical purposes intended for infants below 6 months of age:

- satisfactory, if all the values observed indicate the absence of the bacterium,
- unsatisfactory, if the presence of the bacterium is detected in any of the sample units.

Histamine in fishery products from fish species associated with a high amount of histidine:

- satisfactory, if the following requirements are fulfilled:

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1. the mean value observed is $\leq m$
 2. a maximum of c/n values observed are between m and M
 3. no values observed exceed the limit of M,
- unsatisfactory, if the mean value observed exceeds m or more than c/n values are between m and M or one or more of the values observed are $>M$.

Chapter 2. Process hygiene criteria

2.1. Meat and products thereof

Food category	Micro-organisms	Sampling plan ^a		Limits ^b		Analytical reference method ^c	Stage where the criterion applies	Action in case of unsatisfactory results
		n	c	m	M			
2.1.1.	Aerobic Colony Count of carcasses of cattle, sheep, goats and horses ^d			3,5 log cfu/cm ² daily mean log	5,0 log cfu/cm ² daily mean log	ISO 4833	Carcasses after dressing but before chilling	Improvements in slaughter hygiene and review of process controls
				1,5 log cfu/cm ² daily mean log	2,5 log cfu/cm ² daily mean log	ISO 21528-2	Carcasses after dressing but before chilling	Improvements in slaughter hygiene and review of process controls
2.1.2.	Aerobic Colony Count of carcasses of pigs ^d			4,0 log cfu/cm ² daily mean log	5,0 log cfu/cm ² daily mean log	ISO 4833	Carcasses after dressing but before chilling	Improvements in slaughter hygiene and review of process controls
				2,0 log cfu/cm ² daily mean log	3,0 log cfu/cm ² daily mean log	ISO 21528-2	Carcasses after dressing but before chilling	Improvements in slaughter hygiene and review

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							of process controls
2.1.3.	<i>Salmonella</i> Carcases of cattle, sheep, goats and horses	50 ^e	2 ^f	Absence in the area tested per carcass	EN/ISO 6579	Carcasses after dressing but before chilling	Improvements in slaughter hygiene, review of process controls and of origin of animals
2.1.4.	<i>Salmonella</i> Carcases of pig	50 ^e	5 ^f	Absence in the area tested per carcass	EN/ISO 6579	Carcasses after dressing but before chilling	Improvements in slaughter hygiene and review of process controls, origin of animals and of the biosecurity measures in the farms of origin
2.1.5.	<i>Salmonella</i> Poultry carcasses of broilers and turkeys	50 ^e	7 ^f	Absence in 25 g of a pooled sample of neck skin	EN/ISO 6579	Carcasses after chilling	Improvements in slaughter hygiene and review of process controls, origin of animals and biosecurity measures in the farms of origin

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2.1.6.	Aerobic Mince colony count ^g	5	2	5x10 ⁵ cfu/g	5x10 ⁶ cfu/g	ISO 4833	End of the manufacturing process	Improvements in production hygiene and improvements in selection and/or origin of raw materials
	<i>E.coli</i> ^h	5	2	50 cfu/g	500 cfu/ g	ISO 16649-1 or 2	End of the manufacturing process	Improvements in production hygiene and improvements in selection and/or origin of raw materials
2.1.7.	Aerobic Mechanically separated meat (MSM) ⁱ	5	2	5x10 ⁵ cfu/g	5x10 ⁶ cfu/g	ISO 4833	End of the manufacturing process	Improvements in production hygiene and improvements in selection and/or origin of raw materials
	<i>E.coli</i> ^h	5	2	50 cfu/g	500 cfu/ g	ISO 16649-1 or 2	End of the manufacturing process	Improvements in production hygiene and improvements in selection and/or origin of raw materials

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2.1.8.	<i>E. coli</i> ^h Meat preparations	5	2	500 cfu/ g or cm ²	5 000 cfu/g or cm ²	ISO 16649-1 or 2	End of the manufacturing process	Improvements in production hygiene and improvements in selection and/or origin of raw materials
a	n = number of units comprising the sample; c = number of sample units giving values between m and M.							
b	For points 2.1.3 — 2.1.5 m=M.							
c	The most recent edition of the standard shall be used.							
d	The limits (m and M) apply only to samples taken by the destructive method. The daily mean log is calculated by first taking a log value of each individual test result and then calculating the mean of these log values.							
e	The 50 samples are derived from 10 consecutive sampling sessions in accordance with the sampling rules and frequencies laid down in this Regulation.							
f	The number of samples where the presence of salmonella is detected. The c value is subject to review in order to take into account the progress made in reducing the salmonella prevalence. Member States or regions having low salmonella prevalence may use lower c values even before the review.							
g	This criterion does not apply to minced meat produced at retail level when the shelf-life of the product is less than 24 hours.							
h	<i>E. coli</i> is used here as an indicator of faecal contamination.							
i	These criteria apply to mechanically separated meat (MSM) produced with the techniques referred to in Chapter III, paragraph 3, in section V of Annex III of Regulation (EC) No 853/2004 of the European Parliament and of the Council of 29 April 2004 laying down specific hygiene rules for food of animal origin.							

Interpretation of the test results

The limits given refer to each sample unit tested, excluding testing of carcasses where the limits refer to pooled samples.

The test results demonstrate the microbiological quality of the process tested.

Enterobacteriaceae and aerobic colony count in carcasses of cattle, sheep, goats, horses and pigs:

- satisfactory, if the daily mean log is < m,
- acceptable, if the daily mean log is between m and M,
- unsatisfactory, if the daily mean log is >M.

Salmonella in carcasses:

- satisfactory, if the presence of *Salmonella* is detected in a maximum of c/n samples,
- unsatisfactory, if the presence of *Salmonella* is detected in more than c/n samples.

After each sampling session, the results of the last ten sampling sessions are assessed in order to obtain the n number of samples.

E. coli and aerobic colony count in minced meat, meat preparations and mechanically separated meat (MSM):

- satisfactory, if all the values observed are < m,

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- acceptable, if a maximum of c/n values are between m and M, and the rest of the values observed are < m,
- unsatisfactory, if one or more of the values observed are >M or more than c/n values are between m and M.

2.2. Milk and dairy products

Food category	Micro-organisms	Sampling plan ^a		Limits ^b		Analytical reference method ^c	Stage where the criterion applies	Action in case of unsatisfactory results
		n	c	m	M			
2.2.1.	Enterobacteriaceae Pasteurised milk and other pasteurised liquid dairy products ^d	5	2	<1 cfu/ml	5 cfu/ml	ISO 21528-1	End of the manufacturing process	Check on the efficiency of heat-treatment and prevention of recontamination as well as the quality of raw materials
2.2.2.	<i>E. coli</i> ^e Cheeses made from milk or whey that has undergone heat treatment	5	2	100 cfu/g	1 000 cfu/g	ISO 16649- 1 or 2	At the time during the manufacturing process when the <i>E. coli</i> count is expected to be highest ^f	Improvements in production hygiene and selection of raw materials

a n = number of units comprising the sample; c = number of sample units giving values between m and M.

b For point 2.2.7 m=M.

c The most recent edition of the standard shall be used.

d The criterion does not apply to products intended for further processing in the food industry.

e *E. coli* is used here as an indicator for the level of hygiene.

f For cheeses which are not able to support the growth of *E. coli*, the *E. coli* count is usually the highest at the beginning of the ripening period, and for cheeses which are able to support the growth of *E. coli*, it is normally at the end of the ripening period.

g Excluding cheeses where the manufacturer can demonstrate, to the satisfaction of the competent authorities, that the product does not pose a risk of staphylococcal enterotoxins.

h Only ice creams containing milk ingredients.

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2.2.3.	Coagulase positive staphylococci from raw milk	5	2	10 ⁴ cfu/g	10 ⁵ cfu/g	EN/ISO 6888-2	At the time during the manufacturing process when the number of staphylococci is expected to be highest	Improvements in production hygiene using selection of raw materials. If values >10 ⁵ cfu/g are detected, the cheese batch has to be tested for staphylococcal enterotoxins.
2.2.4.	Coagulase positive staphylococci from milk that has undergone a lower heat treatment than pasteurisation ^g and ripened cheeses made from milk or whey that has undergone pasteurisation or a stronger heat treatment ^g	5	2	100 cfu/g	1 000 cfu/g	EN/ISO 6888-1 or 2		

a n = number of units comprising the sample; c = number of sample units giving values between m and M.

b For point 2.2.7 m=M.

c The most recent edition of the standard shall be used.

d The criterion does not apply to products intended for further processing in the food industry.

e *E. coli* is used here as an indicator for the level of hygiene.

f For cheeses which are not able to support the growth of *E. coli*, the *E. coli* count is usually the highest at the beginning of the ripening period, and for cheeses which are able to support the growth of *E. coli*, it is normally at the end of the ripening period.

g Excluding cheeses where the manufacturer can demonstrate, to the satisfaction of the competent authorities, that the product does not pose a risk of staphylococcal enterotoxins.

h Only ice creams containing milk ingredients.

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2.2.5.	Coagulase positive staphylococci (fresh cheeses) made from milk or whey that has undergone pasteurisation or a stronger heat treatment ^g	5	2	10 cfu/g	100 cfu/g	EN/ISO 6888-1 or 2	End of the manufacturing process	Improvements in production hygiene. If values > 10 ⁵ cfu/g are detected, the cheese batch has to be tested for staphylococcal enterotoxins.
2.2.6.	<i>E. coli</i> ^e Butter and cream made from raw milk or milk that has undergone a lower heat treatment than pasteurisation	5	2	10 cfu/g	100 cfu/g	ISO 16649- 1 or 2	End of the manufacturing process	Improvements in production hygiene and selection of raw materials

a n = number of units comprising the sample; c = number of sample units giving values between m and M.

b For point 2.2.7 m=M.

c The most recent edition of the standard shall be used.

d The criterion does not apply to products intended for further processing in the food industry.

e *E. coli* is used here as an indicator for the level of hygiene.

f For cheeses which are not able to support the growth of *E. coli*, the *E. coli* count is usually the highest at the beginning of the ripening period, and for cheeses which are able to support the growth of *E. coli*, it is normally at the end of the ripening period.

g Excluding cheeses where the manufacturer can demonstrate, to the satisfaction of the competent authorities, that the product does not pose a risk of staphylococcal enterotoxins.

h Only ice creams containing milk ingredients.

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2.2.7.	Enterobacteriaceae	0	10 cfu/g		ISO 21528- 1	End of the manufacturing process	Check on the efficiency of heat treatment and prevention of recontamination
	Milk powder and whey powder ^d						
2.2.7.	Coagulase positive staphylococci	2	10 cfu/g	100 cfu/g	EN/ISO 6888-1 or 2	End of the manufacturing process	Improvements in production hygiene. If values > 10 ⁵ cfu/g are detected, the batch has to be tested for staphylococcal enterotoxins.
2.2.8.	Enterobacteriaceae	2	10 cfu/g	100 cfu/g	ISO 21528- 2	End of the manufacturing process	Improvements in production hygiene
	Ice cream ^h and frozen dairy desserts						
2.2.9.	Enterobacteriaceae	0	Absence in 10 g		ISO 21528- 1	End of the manufacturing process	Improvements in production hygiene to minimise contamination.
	Dried infant formulae and dried dietary foods						

a n = number of units comprising the sample; c = number of sample units giving values between m and M.

b For point 2.2.7 m=M.

c The most recent edition of the standard shall be used.

d The criterion does not apply to products intended for further processing in the food industry.

e *E. coli* is used here as an indicator for the level of hygiene.

f For cheeses which are not able to support the growth of *E. coli*, the *E. coli* count is usually the highest at the beginning of the ripening period, and for cheeses which are able to support the growth of *E. coli*, it is normally at the end of the ripening period.

g Excluding cheeses where the manufacturer can demonstrate, to the satisfaction of the competent authorities, that the product does not pose a risk of staphylococcal enterotoxins.

h Only ice creams containing milk ingredients.

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	for special medical purposes intended for infants below six months of age					If Enterobacteriaceae are detected in any of the sample units, the batch has to be tested for <i>E. sakazakii</i> and <i>Salmonella</i>
a	n = number of units comprising the sample; c = number of sample units giving values between m and M.					
b	For point 2.2.7 m=M.					
c	The most recent edition of the standard shall be used.					
d	The criterion does not apply to products intended for further processing in the food industry.					
e	<i>E. coli</i> is used here as an indicator for the level of hygiene.					
f	For cheeses which are not able to support the growth of <i>E. coli</i> , the <i>E. coli</i> count is usually the highest at the beginning of the ripening period, and for cheeses which are able to support the growth of <i>E. coli</i> , it is normally at the end of the ripening period.					
g	Excluding cheeses where the manufacturer can demonstrate, to the satisfaction of the competent authorities, that the product does not pose a risk of staphylococcal enterotoxins.					
h	Only ice creams containing milk ingredients.					

Interpretation of the test results

The limits given refer to each sample unit tested.

The test results demonstrate the microbiological quality of the process tested.

Enterobacteriaceae in dried infant formulae and dried dietary foods for special medical purposes intended for infants below six months of age:

- satisfactory, if all the values observed indicate the absence of the bacterium,
- unsatisfactory, if the presence of the bacterium is detected in any of the sample units

E. coli, enterobacteriaceae (other food categories) and coagulase-positive staphylococci:

- satisfactory, if all the values observed are < m,
- acceptable, if a maximum of c/n values are between m and M, and the rest of the values observed are < m,
- unsatisfactory, if one or more of the values observed are >M or more than c/n values are between m and M.

2.3. Egg products

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Food category	Micro-organisms	Sampling plan ^a		Limits		Analytical reference method ^b	Stage where the criterion applies	Action in case of unsatisfactory results
		n	c	m	M			
2.3.1.	Enterobacteriaceae Egg products	5	2	10 cfu/g or ml	100 cfu/g or ml	ISO 21528-2	End of the manufacturing process	Checks on the efficiency of the heat treatment and prevention of recontamination

a n = number of units comprising the sample; c = number of sample units giving values between m and M.

b The most recent edition of the standard shall be used.

Interpretation of the test results

The limits given refer to each sample unit tested.

The test results demonstrate the microbiological quality of the process tested.

Enterobacteriaceae in egg products:

- satisfactory, if all the values observed are < m,
- acceptable, if a maximum of c/n values are between m and M, and the rest of the values observed are < m,
- unsatisfactory, if one or more of the values observed are >M or more than c/n values are between m and M.

2.4. Fishery products

Food category	Micro-organisms	Sampling plan ^a		Limits		Analytical reference method ^b	Stage where the criterion applies	Action in case of unsatisfactory results
		n	c	m	M			
2.4.1.	<i>E. coli</i> Shelled and shucked products	5	2	1 cfu/g	10 cfu/g	ISO TS 16649-3	End of the manufacturing process	Improvements in production hygiene
	Coagulase positive staphylococci and molluscan shellfish	5	2	100 cfu/g	1 000 cfu/g	EN/ISO 6888-1 or 2	End of the manufacturing process	Improvements in production hygiene

a n = number of units comprising the sample; c = number of sample units giving values between m and M.

b The most recent edition of the standard shall be used.

Status: This is the original version (as it was originally adopted).

Interpretation of the test results

The limits given refer to each sample unit tested.

The test results demonstrate the microbiological quality of the process tested.

E. coli in shelled and shucked products of cooked crustaceans and molluscan shellfish:

- satisfactory, if all the values observed are $< m$,
- acceptable, if a maximum of c/n values are between m and M , and the rest of the values observed are $\leq m$,
- unsatisfactory, if one or more of the values observed are $> M$ or more than c/n values are between m and M .

Coagulase-positive staphylococci in shelled and cooked crustaceans and molluscan shellfish:

- satisfactory, if all the values observed are $< m$,
- acceptable, if a maximum of c/n values are between m and M , and the rest of the values observed are $< m$,
- unsatisfactory, if one or more of the values observed are $> M$ or more than c/n values are between m and M .

2.5. Vegetables, fruits and products thereof

Food category	Micro-organisms	Sampling plan ^a		Limits		Analytical reference method ^b	Stage where the criterion applies	Action in case of unsatisfactory results
		n	c	m	M			
2.5.1. Pre-cut fruit and vegetables (ready-to-eat)	<i>E. coli</i>	5	2	100 cfu/g	1 000 cfu/g	ISO 16649- 1 or 2	Manufacturing process	Improvements in production hygiene, selection of raw materials
2.5.2. Unpasteurised fruit and vegetable juices (ready-to-eat)	<i>E. coli</i>	5	2	100 cfu/g	1 000 cfu/g	ISO 16649- 1 or 2	Manufacturing process	Improvements in production hygiene, selection of raw materials

a n = number of units comprising the sample; c = number of sample units giving values between m and M.

b The most recent edition of the standard shall be used.

Interpretation of the test results

The limits given refer to each sample unit tested.

The test results demonstrate the microbiological quality of the process tested.

E. coli in pre-cut fruit and vegetables (ready-to-eat) and in unpasteurised fruit and vegetable juices (ready-to-eat):

- satisfactory, if all the values observed are $< m$,
- acceptable, if a maximum of c/n values are between m and M , and the rest of the values observed are $\leq m$,
- unsatisfactory, if one or more of the values observed are $> M$ or more than c/n values are between m and M .

Chapter 3. Rules for sampling and preparation of test samples

3.1. General rules for sampling and preparation of test samples

In the absence of more specific rules on sampling and preparation of test samples, the relevant standards of the ISO (International Organisation for Standardisation) and the guidelines of the Codex Alimentarius shall be used as reference methods.

3.2. Bacteriological sampling in slaughterhouses and at premises producing minced meat and meat preparations

Sampling rules for carcasses of cattle, pigs, sheep, goats and horses

The destructive and non-destructive sampling methods, the selection of the sampling sites and the rules for storage and transport of samples are described in standard ISO 17604.

Five carcasses shall be sampled at random during each sampling session. Sample sites should be selected taking into account the slaughter technology used in each plant.

When sampling for analyses of enterobacteriaceae and aerobic colony counts, four sites of each carcass shall be sampled. Four tissue samples representing a total of 20 cm² shall be obtained by the destructive method. When using the non-destructive method for this purpose, the sampling area shall cover a minimum of 100 cm² (50 cm² for small ruminant carcasses) per sampling site.

When sampling for *Salmonella* analyses, an abrasive sponge sampling method shall be used. The sampling area shall cover a minimum of 100 cm² per site selected.

When samples are taken from the different sampling sites on the carcass, they shall be pooled before examination.

Sampling rules for poultry carcasses

For the *Salmonella* analyses, a minimum of 15 carcasses shall be sampled at random during each sampling session and after chilling. A piece of approximately 10 g from neck skin shall be obtained from each carcass. On each occasion the neck skin samples from three carcasses shall be pooled before examination in order to form 5 x 25 g final samples.

Guidelines for sampling

More detailed guidelines on the sampling of carcasses, in particular concerning the sampling sites, may be included in the guides to good practice referred to in Article 7 of Regulation (EC) No 852/2004.

Sampling frequencies for carcasses, minced meat, meat preparations and mechanically separated meat

However, when justified on the basis of a risk analysis and consequently authorised by the competent authority, small slaughterhouses and establishments producing minced meat and meat preparations in small quantities may be exempted from these sampling frequencies.

Status: This is the original version (as it was originally adopted).

In the case of sampling for *Salmonella* analyses of minced meat, meat preparations and carcasses, the frequency can be reduced to fortnightly if satisfactory results have been obtained for 30 consecutive weeks. The salmonella sampling frequency may also be reduced if there is a national or regional salmonella control programme in place and if this programme includes testing that replaces the described sampling. The sampling frequency may be further reduced if the national or regional salmonella control programme demonstrates that the salmonella prevalence is low in animals purchased by the slaughterhouse.

As regards the sampling of minced meat and meat preparations for *E. coli* and aerobic colony count analyses and the sampling of carcasses for enterobacteriaceae and aerobic colony count analyses, the frequency may be reduced to fortnightly testing if satisfactory results are obtained for six consecutive weeks.

The food business operators of slaughterhouses or establishments producing minced meat, meat preparations or mechanically separated meat shall take samples for microbiological analysis at least once a week. The day of sampling shall be changed each week to ensure that each day of the week is covered.

- (1) The test results can be used also for demonstrating the effectiveness of the HACCP or good hygiene procedure of the process.