

Audit checklist Checklist Slaughtering/Deboning (Spotaudit)

Audit details				
Scheme participant				
QS locations audited				
Additional location information, e.g. coordinators or identification number				
Name of contact				
Spotaudit	Х			
Random sample audit				
Audit of special purpose				
Parallel audit				
Date of audit (from)			Date of audit (until)	
Start of audit (hh:mm)			Ende of audit (hh:mm)	
Audit duration (hh:mm)				
Combined audit (norm/standard/programme)				
Certification body				
First name/surename of auditor				
Repeated D evaluation/general K.O.		Remark repeated D evaluation/general K.O.		
Comments				
Preliminary audit results			Number of agreed corractions	rective
				-
Place, date		Signature/s of a	uditor/s	
I hereby confirm the data concerr I have received a copy of the aud	_			ons report.
Place, date		Signature of per	son responsible	



Company details - Slaughtering/deboning

Name of company	
Street and house number	
Postal code and town	
Telephone/fax number	
Email address	
Registered production scope no.	
QS location number	
QS identification number	
Name of person responsible	

Scope - Slaughtering/deboning

Product	ion scope	Production number
	Slaughtering beef, veal, pork	31
	Deboning beef, veal, pork	32
	Slaughtering poultry	34
	Deboning poultry	35
Tonnag	e per year	



Company									Date	
Require ment no.	Factor	Filter ¹		Criterion/ requirement	A	В	С	D/ K.O.	E	Comments/corrective action number
complia the asse	nce v	with tent.	the QS r # = In c	the evidence or measu equirement must be do ase of a nonconformity ays (only valid for prod	cum the	ent cori	ed, ı ecti	regard ve ac	dles: tion	s of the outcome of for this criterion has
2	Gen	eral r	equireme	nts						
2.1	Gen	eral s	cheme re	quirements						
2.1.1	1			General Business data					Х	
2.1.2	1			Incident and crisis management					Х	
2.1.3	1			Disaster concept					Х	
2.1.4	1			Food safety culture *					Х	
2.1.5	1			Appointing service providers *					Х	
2.2	Self	-asse:	ssment a	nd HACCP						
2.2.1	1		D=K.O.	Conducting self- assessment					Х	
2.2.2	1			Listeria monitoring *					Х	
2.2.3	1			Document handling					Х	
2.2.4	1		D=K.O.	HACCP- Konzept/Managementsyste me für					Х	
2.2.5	1			HACCP-Team					Х	
	-									



Require ment no.	Factor	Filter ¹		Criterion/ requirement	A	В	С	D/ K.O.	E	Comments/corrective action number
2.2.6	1			Product description					Х	
2.2.7	1			Flow chart					Х	
2.2.8	1			Hazard analysis					Х	
2.2.9	1			Critical Control Point (CCP)					Х	
2.2.10	1			Limit values for CCP					X	
2.2.11	1			Monitoring and verification of limit values for CCP					Х	
2.2.12	1			Corrective actions for CCP					Х	
2.2.13	1			Responsibilities					Х	
2.2.14	1			Documentation					Х	
2.2.15	1			HACCP verification					Х	
2.3	Goo	d man	ufacturin	g and hygiene practice						
2.3.1	1			Water quality *					Х	
2.3.2	1			Development of cleaning and disinfection plans					Х	
2.3.3	1		D=K.O.	Microbiological control of cleaning and disinfection measures					Х	
2.3.4 SPOT	1			Foreign matter management *						
2.3.5	1			Production release *					Х	
			1	1			•			



Require ment no.	Factor	Filter ¹		Criterion/ requirement	A	В	С	D/ K.O.	E	Comments/corrective action number
2.3.6	1			Pest control					Χ	
2.3.7	1			Maintenance and repair					Х	
2.3.8	1			Monitoring of Test Equipment					Х	
2.3.9 SPOT	1		D=K.O.	Contamination *						
2.3.10	1			Allergen Management					Х	
2.3.11	1			Species-specific product separation					X	
2.5				al condition						
2.6	Gro	und cl	earance							
2.7	Sta	ff								
2.7.1 SPOT	1			General rules of conduct and Staff hygiene						
	-									
2.7.2 SPOT	1			Premises and Access Regulations						
	1									
SPOT 2.7.3			D=K.O.	Regulations Staff rooms and sanitary						
SPOT 2.7.3 SPOT 2.7.4	1	ning (D=K.O.	Regulations Staff rooms and sanitary facilities						



			ı					ı		
Require ment no.	Factor	Filter ¹		Criterion/ requirement	A	В	С	D/ K.O.	E	Comments/corrective action number
2.8.2	1			Information on the QS scheme					Х	
3	Aniı	mal w	elfare							
3.1	Gen	eral r	equireme	nts						
3.1.1 SPOT	1		D=K.O.	Animal welfare officer						
3.1.2	1			Standard work instructions					Х	
3.1.3	1			Employee competence *					Х	
3.1.4 SPOT	1		D=K.O.	Livestock handling						
3.2	Aniı	mal w	elfare in t	he shed/sty area				 		
3.2.1 SPOT	1			Water dispensers feeding and bedding						
3.2.2 SPOT	1			Climate conditions						
3.2.3 SPOT	1			Sprinkler system						
3.2.4 SPOT	1			Crates allocation						
3.3	Aniı	mal W	elfare in t	the stunning area						
3.3.1 SPOT	1			Stunning system *						
3.3.2 SPOT	1			Driving livestock to the stunning area *						
3.3.3 SPOT	1		D=K.O.	Effective stunning *						



Require ment no.	Factor	Filter ¹		Criterion/ requirement	A	В	С	D/ K.O.	E	Comments/corrective action number
3.3.4 SPOT	1			Re-stunning *						
4	Slau	ıghter	requiren	nents						
4.1	Live	stock	transpor	t monitoring - transport p	racti	ce				
4.1.1	1		D=K.O.	Verification animal transporter					Х	
4.1.2 SPOT	1			Delivery						
4.1.3 SPOT	1		D=K.O.	Verifying the indication of origin and delivery authorization of QS						
4.2 Ramp area, shed/shy, waiting area										
4.2.1 SPOT	1			Unloading facilities						
4.2.2 SPOT	1			Separation from animals						
4.2.3	1			Technical/structural condition					Х	
4.2.4 SPOT	1			Premises, facility and device hygiene						
4.3	Slau	ighter	process		Į.					
4.3.1 SPOT	1			Shackling and hoisting						
4.3.2 SPOT	1			Bleeding						
4.3.3 SPOT	1			Skinning/bristle removal/plucking						
4.3.4 SPOT	1			Removal of stomach and chest organs						



Require ment no.	Factor	Filter1		Criterion/ requirement	A	В	С	D/ K.O.	E	Comments/corrective action number
4.3.5 SPOT	1			Carcass splitting						
4.3.6 SPOT	1		D=K.O.	Sluice option						
4.3.7 SPOT	1			Post-processing line						
4.3.8 SPOT	1			Technical/structural condition						
4.3.9 SPOT	1			Premises, facility and device hygiene						
4.3.10 SPOT	1		D=K.O.	Organisation and workflows						
4.3.11 SPOT	1			Blade hygiene						
4.3.12 SPOT	1			Climate conditions						
4.3.13	1		D=K.O.	Diagnostic data pig *					Х	
4.3.14	1		D=K.O.	Diagnostic data cattle *					Х	
4.3.15	1		D=K.O.	Diagnostic data poultry *					Х	
4.3.16	1		D=K.O.	Salmonella monitoring *					X	
4.3.17	1			Logistical slaugthering of salmonella-positive herds (poultry)					Х	
4.3.18	1			Turkey slaughtering: participation in PAI monitoring					Х	
4.3.19	1			Taint detection					Х	
4.4	4 Cold storage (carcasses)									



Require ment no.	Factor	Filter ¹		Criterion/ requirement	A	В	С	D/ K.O.	E	Comments/corrective action number
4.4.1 SPOT	1			Technical/structural condition						
4.4.2 SPOT	1			Premises, facility and device hygiene						
4.4.3 SPOT	1			Ground clearance						
4.4.4	1			Storage management					Х	
4.4.5	1		D=K.O.	Temperature recording and monitoring after slaughter					Х	
4.4.6 SPOT	1			Quartering cattle						
5	Req	uirem	ents for o	leboning						
5.1	Deb	oning								
5.1.1 SPOT	1			Technical/structural condition						
5.1.2 SPOT	1			Premises, facility and device hygiene						
5.1.3 SPOT	1			Ground clearance						
5.1.4 SPOT	1		D=K.O.	Organisation and workflow						
5.1.5 SPOT	1			Handling of non- conforming products						
5.1.6 SPOT	1		D=K.O.	Temperature recording and monitoring *						
5.2	Cut	ting, p	ortioning	and minced meat product	ion					
5.2.1 SPOT	1			Technical/structural condition						



Tequent no. The second	ene arance	АВ	C C	D/ K.O.	E	Comments/corrective action number
SPOT device hygi 5.2.3 1 Ground clea SPOT D=K.O. Organistaio	arance n and					
SPOT D=K.O. Organistaio	n and					
SPOT workflows	e recording and					
5.2.5 1 D=K.O. Temperatur monitoring	e recording and				Х	
5.3 Labelling and packaging					<u> </u>	
5.3.1 1 Technical/st	ructural					
5.3.2 1 Premises, for device hygi	-					
5.3.3 1 D=K.O. Packaging r	material *				Х	
5.3.4 1 D=K.O. Final production	ct inspection					
5.3.5 1 D=K.O. Product lab	elling				Х	
5.3.6 1 D=K.O. Recipes/spe	ecifications *				Х	
5.3.7 1 D=K.O. Final production plans *	ct sampling				Х	
5.4 Meat cold storage (packaged	goods)					
5.4.1 1 Technical/st condition	ructural					
5.4.2 1 Premises, for device hygi						
5.4.3 1 Ground clea	arance					



Require ment no.	Factor	Filter ¹		Criterion/ requirement	A	В	С	D/ K.O.	E	Comments/corrective action number
5.4.4	1			Storage management					Х	
5.4.5	1		D=K.O.	Temperature recording and monitoring					Х	
5.5	5.5 Deep-freeze facility									
5.5.1 SPOT	1			Technical/structural condition						
5.5.2 SPOT	1			Premises, facility and device hygiene						
5.5.3 SPOT	1			Ground clearance						
5.5.4	1			Storage management					Х	
5.5.5	1		D=K.O.	Temperature recording and monitoring					Х	
6	Add	itiona	l producti	on departements and faci	lities	5				
6.1	Clea	aning	rooms and	d material storage						
6.1.1 SPOT	1			Cleaning rooms						
6.1.2 SPOT	1			Packaging material storage						
6.1.3 SPOT	1			Cleaning product and disinfection storage						
6.1.4 SPOT	1		D=K.O.	Spice storage *						
6.2	Dis	oosal				ı				
6.2.1	1			Disposal logistics					Х	



Require ment no.	Factor	Filter ¹		Criterion/ requirement	A	В	С	D/ K.O.	E	Comments/corrective action number	
6.2.2 SPOT	1			Disposal area							
6.2.3 SPOT	1		D=K.O.	Slaughter by-products and risk material							
6.3	Vehicle fleet										
6.3.1	1			Transport vehicle washing facilities					Х		
6.3.2	1			Cleaning and disinfection					X		
6.3.3 SPOT	1			Temperature monitoring system							
7	Pur	chase,	traceabi	lity, labelling, use of the c	ertif	icati	on m	ark ar	nd go	oods separation	
7.1	Inc	oming	and outg	oing goods							
7.1.1 SPOT	1			Technical/structural condition							
7.1.2 SPOT	1			Premises, facility and device hygiene							
7.1.3 SPOT	1			Ground clearance							
7.1.4 SPOT	1		D=K.O.	Incoming goods inspection							
7.1.5 SPOT	1		D=K.O.	Outgoing goods inspection							
7.1.6	1		D=K.O.	Returns management					Х		
7.1.7	1			Claims management					Х		
7.2	Lab	elling	and use (of certification mark			<u> </u>				



Require ment no.	Factor	Filter¹		Criterion/ requirement	A	В	С	D/ K.O.	E	Comments/corrective action number	
7.2.1 SPOT	1		D=K.O.	Labelling of marketed QS goods *							
7.2.2	1			Use of QS certification mark *					Х		
7.3	Traceability and origin of goods										
7.3.1	1		D=K.O.	Traceability method					Х		
7.3.2	1		D=K.O.	Traceability check *					Х		
7.3.3	1		D=K.O.	Quantity comparison					Х		
7.3.4 SPOT	1		D=K.O.	Eligibility of delivery check							
7.4	Goo	ds sep	paration								
7.4.1 SPOT	1		D=K.O.	Separation and identification of QS produce/non-QS produce *							
D 1	Ad on Convenience										
D 2.1	Gen	eral s	cheme re	quirements							
D 2.1.1	1			Use of the QS certification mark					Х		
D 2.2	Good Manufacturing and hygiene practises										
D 2.2.1	1		D=K.O.	Recipes/Spcifications					Х		
D 2.3	Tec	hnical	/structura	al condition							
D 2.4	Pre	mises,	facility a	nd device hygiene							



		1	ı					1			
Require ment no.	Factor	Filter ¹		Criterion/ requirement	A	В	С	D/ K.O.	E	Comments/corrective action number	
D 2.5	Ground Clearence										
D 3.1	Requirements for the production process										
D 3.1.1	1 Best-before date/use-by date										
D 4.1	Silo	stora	ge								
D 4.1.1 SPOT	1			Silo storage					Х		
D 4.1.2 SPOT	1			Technical/structural condition					Х		
D 4.1.3 SPOT	1			Premises, facility and device hygiene			Х				
D 5.1	Tan	k stor	age								
D 5.1.1 SPOT	1			Tank storage					Х		
D 5.1.2 SPOT	1 Technical/structural condition					Х					
D 5.1.3 SPOT	1			Premises, facility and device hygiene					Х		
D 6.1	Pre	paratio	on and Pro	ocessing							
D 6.1.1 SPOT	1			Technical/structural condition					Х		
D 6.1.2 SPOT	1			Premises, facility and device hygiene					Х		
D 6.1.3 SPOT	1			Ground clearence					Х		
D 6.1.4 SPOT	1			. Organistaion and workflows					Х		



Require ment no.	Factor	Filter ¹		Criterion/ requirement	A	В	С	D/ K.O.	E	Comments/corrective action number		
D 7.1												
D 7.1.1 SPOT	1			Technical/structural condition					Х			
D 7.1.2 SPOT	1			Premises, facility and device hygiene					Х			
D 7.1.3 SPOT	1			Ground clearence					Х			
D 7.1.4 SPOT	1			Ordnung und Organisation								
D 8.1	Furt	her pr	ocessing									
D 8.1.1 SPOT	1			Technical/structural condition					Х			
D 8.1.2 SPOT	1			Premises, facility and device hygiene					Х			
D 8.1.3 SPOT	1			Ground clearence					Х			
D 8.1.4 SPOT	1			Organistaion and workflows					Х			
I. VLOG-Additional Module												
I. 1	Req	uirem	ent (only	relevant for locations reg	ister	ed fo	or VL	.OG-A	dditi	onal Module)		
I. 1.1	0			Requirement "Ohne Gentechnik"					Х			



Company	Date											
Calculation of a	udit res	sult										
1. Balance of sub												
Calculation					Α	В	С	D	E			
(1) Number of eval	uations											
Sum of evaluation	ns (exclı	uding E evaluat	tions)									
2. Calculation of	the prop	ortion of C and	D evaluations*									
Proportion	of C eva	luations			(Nu	umber of C ev	aluations / sum o	of evaluations)*	*100			
Proportion	of D eva	luations			(Nu	umber of D ev	aluations / sum o	of evaluations)*	*100			
	on of Ca aluations					Propo	rtion of C + propo	ortion of D				
3. Preliminary au	dit resul	t										
			Percentage of C evaluations		tage of uations	C	ntage of C+D uations	Audit	result			
			max. 5,0%	0,0	0%			QS-Sta	atus I*			
*Status I: If the 5 % is exceeded, status I w			max. 10,0%	max.	3,0%	max. 10%		QS-Status II**				
be assigned if there is	only one Status		max. 20%	max.	10% max. 2		ex. 20% QS-		QS-Status III			
regard to the proporti evaluations is exceede II is assigned if only o evaluation exists and evaluation	on of D ed, status one D	Percentages exceeded			Audit not passed.							
Number of K.O.		K.O.			Audit n	Audit not passed.						
	,	General K.O./ repeated D	Audit not passed.									



Company:	Date:
----------	-------

Corrective actions report

I hereby confirm that the following corrective actions were agreed upon between me and the auditor.

The certification body is to be informed no later than the expiry of the deadline set out in the action plan about the implementation of a corrective action. Note: The correction deadline is a maximum of 28 days for all FIAS requirements and the documentation requirements: 2.1.1, 2.1.2, 3.4.1 und 3.9.5 (only valid for production!)

Place, date

Signature/s of auditor/s

Signature of person responsible

Requirement No.

Requirement No.

Description of nonconformity

Agreed corrective actions

Scope

Deadline for correction



Date:

Review of the implementation of corrective actions									
Place, date			Signature/s of auditor/s						
Serial no.	Implemented	Not implemented	Comments (if any)	Date					
1									

Company: