

### iMIS Food Compliance demo

# the knowledge, content, infrastructure and processes for food safety, legislation and certification

#### Do you ever wonder if you...

- Have a QA manager next month.
- ✓ Will have no surprises in an unannounced audit.
- Comply with the correct legislation.
- Comply with the standards and customer requirements.
- Can double the productivity of the QA department.

# You can manage Food Safety yourself with iMIS Food!







### About QAssurance and Training

#### **QAssurance Roadmap**

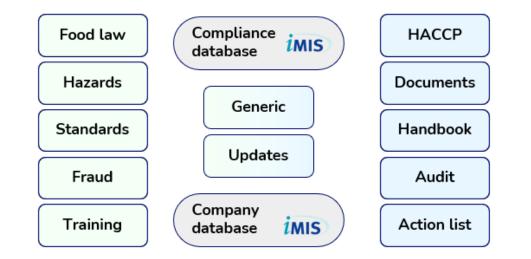
A Journey of Growth and Innovation in Food Safety and Compliance



		1993	1993-1997	1997-2003	2003-2006	2006-2021	2022	2023	
	COMPANY	University TUE	Family Lab Consultant	Unilever	QA Interim	QAssurance	QAssurance	QAssurance	
imis	IMIS FOOD DEVELOPMENT	HACCP System Development	Student & QA Consultant	Learning Organization	QMS Approach Software	QMS License Model	English, German, French Version	iMIS Food Compliance-Ready for Global Roll-out	
F	MILESTONES	HACCP Thesis Cees	Food Safety Authority Approaved (1996)	Global Food Supply Chain Group	First GFSI Certification (2004)	Dutch HACCP Organization >1000 GFSI Audits	GBA Group (DACH Partner), Van Hall University	Agrea, WUR, FoodChain ID	
\$	COUNTRIES	Netherlands	Netherlands	Netherlands, UK, India	Netherlands, UK, India	Netherlands, Belgium, Rwanda (2021)	African Countries (online), Jordan	Philippines, Nigeria, Thailand, Saudi Arabia	



#### iMIS Food Compliance



#### **iMIS Food offers**

Plan a Demo

- The knowledge, content, infrastructure, systems and real-time updates for Food Safety Compliance;
- Always compliant with EU legislation and standards;
- Direct access to our Food Safety experts;
- The system also works offline;
- Dutch, English, German and French content.







#### Food Safety Compliance management

Management of business performance, demand and supply relations, legislation and certification with regard to Food Safety, taking into specifications, quality activities, traceability and assessment.

	Specifications	Quality activities	Traceability	Assessment
	Requesting, drawing up, issuing and securing specifications regarding raw materials, semi-finished products, processes and end products, with which legalislation, quality standards and customer requirements are met.	Drawing up, complying with and guaranteeing product and process parameters through procedures, job descriptions and responsibilities with which legalislation, quality standards and customer requirements are met.	Registering all information flows and related actions regarding raw materials, semi-finished products, processes and end products, with which legalislation, quality standards and customer requirements are met (transparency and consumer intimacy).	Testing whether the product and process parameters and the related procedures, job descriptions and responsibilities meet legislation, the quality standards and customer requirements that are set.
Quality Standards				
Certification	Standard requirements	Operational framework	Test, certification body informed	HACCP, TACCP, VACCP, standard based practice
Lustomers & Consumers				
Demand Relationships	Product, process requirements	Demand Information Center	Products, specifications	Customer satisfaction, consumer needs
Food Company				
Food Company	Product, process and people requirements	Training, support, procedures, quality documents and database	Ingredients, semi-products, final products	Business System
i Suppliers				
Supply Relationships	Product, process and people requirements	Supply Information Center	Raw materials, services, specifications	Supplier selection and performance
✓ Legislation				
Legislation	Legal requirements	HACCP and prerequisite program	Food Safety Authority informed	HACCP, legal based practice





#### Explanation of the case cockpit

- The implementation steps are:
  - 1. Studies, like HACCP and Food legislation
  - 2. Quality System, setting up iMIS
  - 3. Registrations, working with iMIS
  - 4. Reports, making the yearly reports
  - 5. Audit & Support, support by advisers
- You can use next slide as navigator: the customer case cockpit.
- Click on the buttons to go to the next slides for clarification.

Cloud	Food Law	)	Audit rounds		
Server	HACCP	QMS specific	Action list	GFSI reports	GFSI audit
Faad Law Compliance IMS HACCP Matante Matante Documents Standards Generic Handbook	VACCP	QMS generic	Report		Helpdesk
Training Company Justs Adding	ТАССР	)	Documents		
Implementation steps	1. Studies	2. Quality System	3. Registrations	4. Reports	5. Audit & Support
Hazards & Fraud	x			х	х
Legislation	х	х		x	х
Standards	х			х	х
Generic	x	x		х	х
<u>Generic</u> Handbook	х	x x	x	x x	x x
	x		x x		



- With the <case> button on the other slides you can go back to the navigator.
- the relevant iMIS modules are specified for each step with an arrow.

Case	
iMIS Food modules	
Hazards & Fraud	
Legislation	
Standards	
Generic	
Handbook	
Audit & Ticket	
Documents	

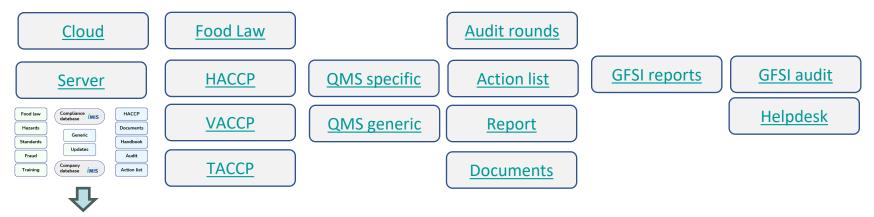
*Normally every step needs around 3-4 days of advice.* 

An implementation is normally executed within 3-6 months.





# Case cockpit: iMIS Food implementation

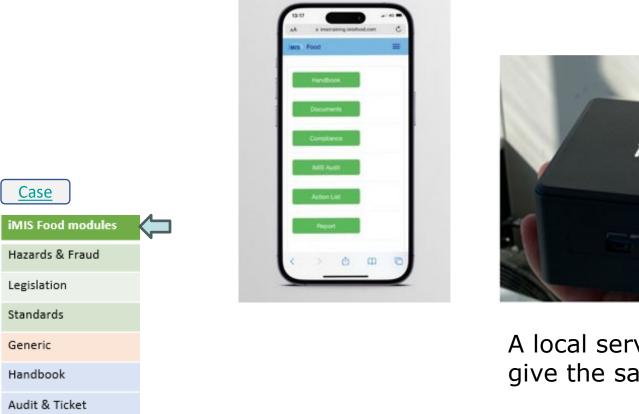


Implementation steps	1. Studies	2. Quality System	3. Registrations	4. Reports	5. Audit & Support
Hazards & Fraud	х			Х	Х
Legislation	х	Х		Х	Х
<u>Standards</u>	х			Х	Х
<u>Generic</u>	х	Х		х	х
Handbook		Х	Х	Х	Х
Audit & Action List			х	х	х
<u>Documents</u>			Х	Х	Х





#### Cloud: mobile functions





A local server with wifi will give the same functionality







#### Food Law

Case

Legislation

Standards

Generic

Handbook

Audit & Ticket

Documents

iMIS Food modules

Hazards & Fraud

#### Legislation overview

	Regulation Directive	Description	Animal by products	Animal feed		Hospitality     and Craft     production		G Foodstuffs     plant based		Composite	Second For specific groups	Special Food and	Supplements	🚱 Dairy
		TSE Regulation	Х	Х										
	1069/2009	The legal framework for control and supervision of animal products	Х	Х	Х									Х
		Implementing regulation	Х	Х	Х									Х
	178/2002	General principles and requirements of food law establishing a European Food		X	X	Х	Х	Х	X	Х	X	Х	х	Х
	-legislation-per-pro animal-by-products													
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	ish-and-seafood													
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### HACCP

	Study H	ACCP Ar	nalysis o	of Process	ses										
	Process step	Hazard	Kind of hazard	Cause	Potential effect (see explanation)	occurrence	likelihood x potential	Control measure(s)	After implementation of the basic conditions program (PRP), the normal process, and taking into account the intended and expected use of the product in the chain after us, is here still an unacceptable risk to the safety or quality of the product?	eliminated or sufficiently reduced by a (combination of) specific control	measure have critical limit value(s)? Is there a legal requirement?	CCP: Critical control point or OPRP or PRP. If failure of the control measure leads to a high risk for food safety and there is a critical value (limit) for the process and it is a single control measure (no combination that can still absorb the failure), then it is a CCP.	correct and detect a deviation	deviations (severity	No. substantiation
	General														Prevention of cross-contamination (hygiene, allergens) and contamination with foreign bodies must be controlled through the PRP. Among other things, through the cleaning plan, allergen management, hygiene procedures, maintenance, et
	Purchasing	See raw materials analysis	See raw materials analysis	See raw materials analysis				Approval and rating of suppliers. Verification. Specifications. Contracts, agreemensts.							Purchasing at approved suppliers.
	Receiving frozen products (fruit maybe in the future)	Growth of pathogens	Microbiological	Wrong conditions during transport	2	1	2	Control of incoming goods. Label, best before date, temperature, other conditions (e.g. hygiene). Selection and assessment of suppliers and purchasing on specification. Product temperature control	No		Legal	PRP	High	Low	The transport temperature of frozen products is regulated by legitation. This must be -18° C and may rise to -15° C for short period of time. However, this has nothing to do with public health hazards. Other conditions are checked by defa at the moment of reaeving the goods. It is only when defrosting that a risk to food safety arises.
	Receiving vegetables (carrots, zuchini, potatoes)	Growth of micro organisms or other spoilage.		Wrong conditions during storage before transport		2	4	Control of incoming goods. Label, best before date, temperature, other conditions (e.g., hygien). Selection and assessment of suppliers and purchasing on specification. Visual check.	No			PRP	High	Low	The product is stored under 4-10°C. In the event of a short-term rise in temperature, there is no immediate hazard. A risk arises if the ambient temperature rises above the maximum value for a longer period of time and the product will deteriorist (spollage). With raw materials that are still heated in the process or before consumption, risk is low, Controlling the temperature until delivery is a responsibility of the supplier. Transporting time is within a day. It is unprocessed potato, carrot and zucohini.
<u>Case</u>	Receiving ambient raw materials like sunflower oil and seasonings.	No specific hazard					-	Control of incoming goods. Label, best before date, temperature, other conditions (e.g. hygiene), Selection and	No			PRP	High	Low	
IS Food modules	Receiving packaging materials and othe	Had	сср	Stu	Idie	Э		assessment of suppliers and purchasing on specification. Ingangscontrole. Selectie en beoordeling van leveranciers en inkoop op specificatie.	Nee			PRP	High	Low	Zie raw materials analysis, A specification and DoC is needed.
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#### VACCP

Case

#### FOR IDENTIFIED RISKS: &

- Analysis certificates.
- 2. Mass balance testing at the supplier for critical ingredients.
- 3. Audits on the chain.
- 4. Track changes in the chain, eliminate links if possible.
- 5. Check middlemen, etc.
- 6. Choose any alternatives with lower risk.
- 7. Other origin (country).
- 8. Improve relationships with suppliers.
- 9. Adjust purchasing policy; prevent spot markets where traceability is difficult.

#### ANNEX RISK INVENTORY RAW MATERIALS

Rating (PRN)	Probability of occurrence (O)	Probability of detection (D)	Profitability (P)
1	Very unlikely or not	Definitely	Very low
2	Unlikely or small	High	Low
3	Medium or significant	Fairly unlikely	Medium or significant
4	High	Unlikely	High
5	Very high	Very unlikely	Very high

🖽 Edit

PRN = O x D x P, action limit 20. In this appendix, this risk assessment is at ingredient level within raw materials.

The probability of occurrence is currently estimated at 1 or 2 for each supplier, because we act on the basis of historical data with our own suppliers. With new suppliers, the PRN score will probably exceed 20 earlier, which requires extra action.

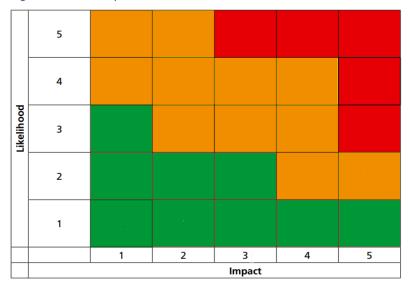
Raw material	Occurence	Detection	Profitability	PRN	Conclusion
Honey and bee products, royal jelly	2	2	4	16	No additional measure needed
Packaging	1	2	1	2	No additional measure needed

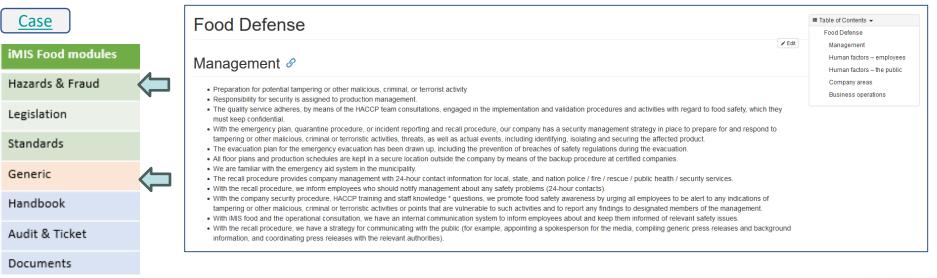
iMIS Food modules	
	A vulnerability assessment is executed on the basis of this information, on all groups of foodstuff raw materials in order to assess the risk of counterfeiting or substituting. This takes into account:
Hazards & Fraud	
Legislation	Refinement of routine testing to identify counterfeit resources;     Nature of the raw material.
	This vulnerability assessment is formally assessed at least annually in order to take into account and process modified economic conditions and market information that could change the possible risks.
Standards	If raw materials are identified in this vulnerability assessment as being at special risk of counterfeiting or substitution, appropriate warranty and / or testing processes will be put into place to reduce the risk.
Standards	If products are labelled or there are claims made on the end packaging that depend on the status of the raw material, such as:
Generic	Specific origin or source;     Specific origin or source;     Status of assurance (for example, Global GAP);     Genetically modified organism status (GMO);
Handbook	Identify retention;     Specific name trademark ingredients.     then the status of each batch of raw material will be checked.
Audit & Ticket	As an organization, we record putchasing data, traceability of use of raw materials and end product packaging with claims. As an organization we are obligated to frequently perform a documented mass balance test with regard to claims. This is carried out and recorded at least every 6 months (in case of the absence of a system certificate) that guarantees product authenticity.
	For claims related to the methods of production (for example organic, Beter Leven, halal, kosher), we as an organization must have the necessary certification to be able to make this claim.
Documents	The process flow for the product of products with a claim is documented and possible areas for contamination or loss of identity identified. Appropriate controls are established to ensure product integrity.
	Associated files
	Food Fraud Vulnerability Assessment.
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#### TACCP

Figure A.5 – Threat prioritization

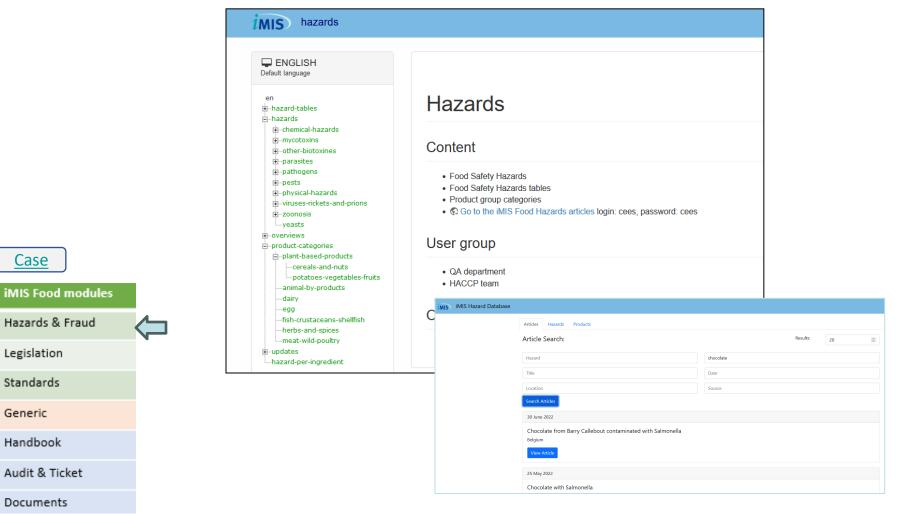








#### **iMIS** Food Hazards





Case

Legislation

Standards

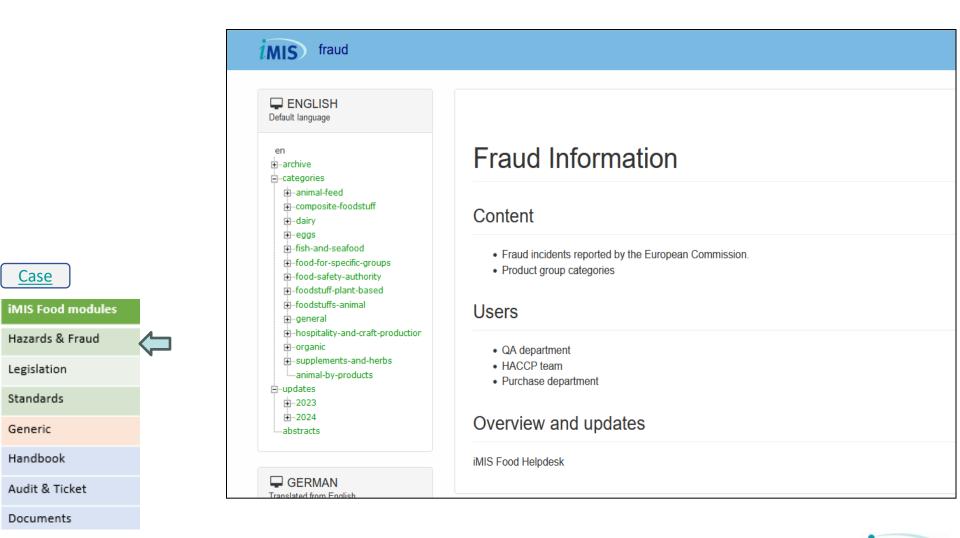
Generic

Handbook

Documents



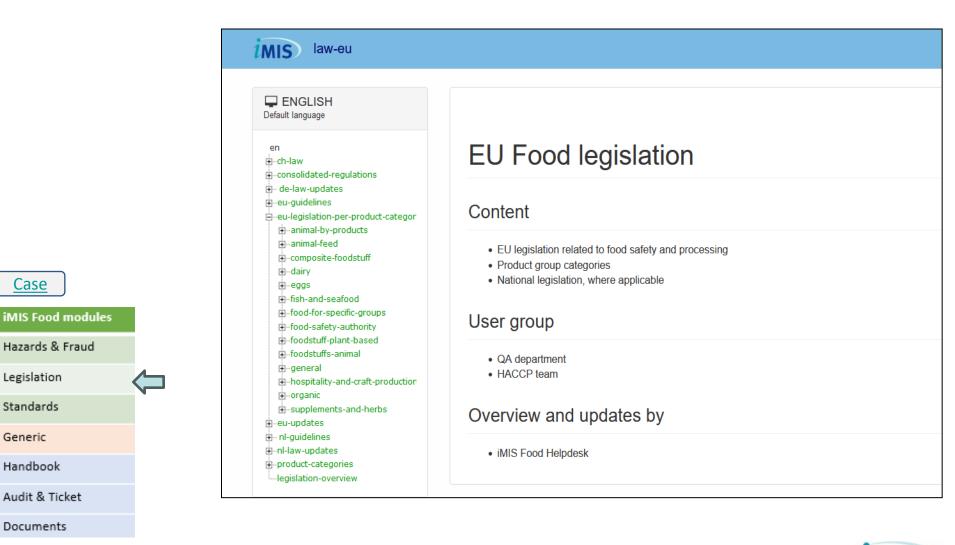
### iMIS Food Fraud





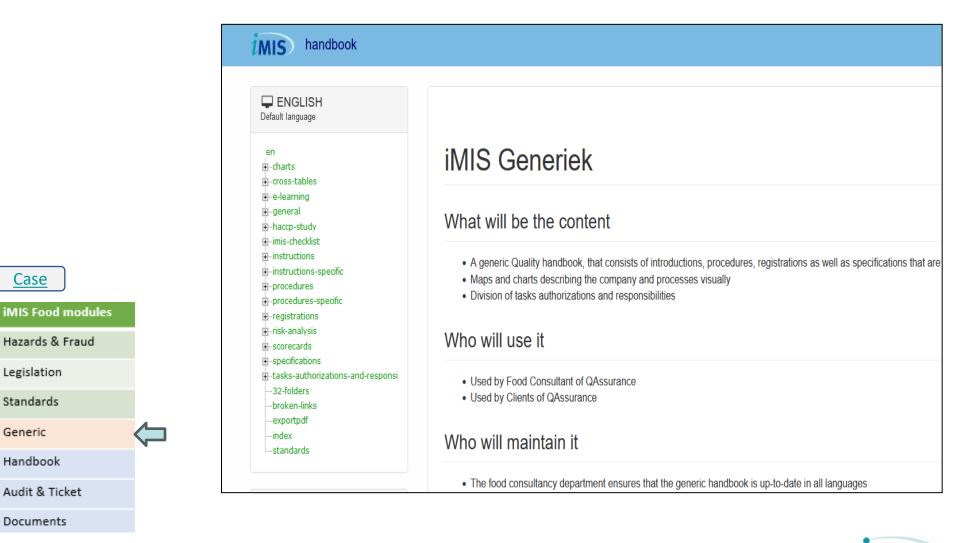


#### iMIS Food Legislation





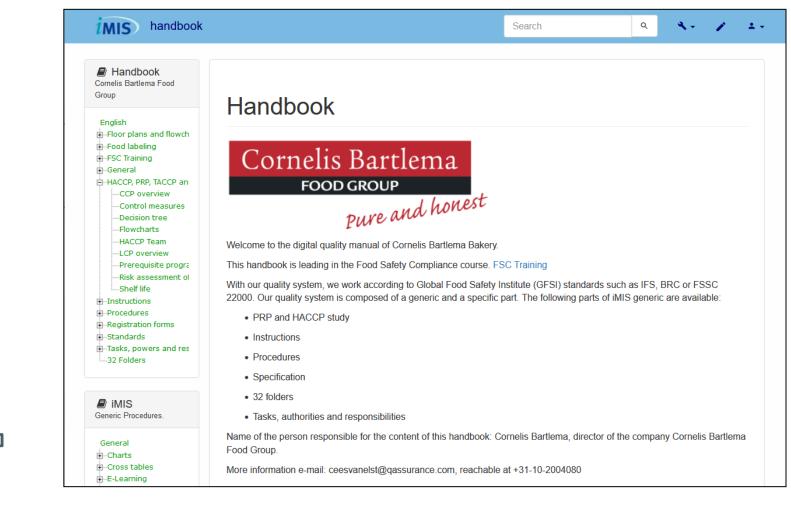
#### iMIS Food Generic







#### iMIS Food Handbook



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Audit & Ticket

Documents

iMIS Food modules

Hazards & Fraud



### iMIS Food Audit & Report

Haccp team meet	ing [11]				
<u>Detail</u> ►		Action Latt			
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Entire team	<b>⊘</b> 1	imis Audit Reports	Sear	ch Q	y.
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Haccp team meeting takes place on time	<b>⊘</b> 1	Audit Poporto			
		Audit Reports			
Reception			Search/Filter		
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No change in products	<b>⊘</b> 1	any v			
No change in machines	⊘ 1	v any: haccp			
No new employees	⊘ 1		Apply filters		
Sufficient knowledge of staff	<b>⊘</b> 1	← Previous page	1		Next
		21 96% Haccp team meeting [21]		29-5-2024 10:41:00 Aysha	
		11 99% Haccp team meeting [11]		10-3-2024 9:41:00 Cees	
		← Previous page	1		Next

Documents

Audit & Ticket

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Standards

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Handbook

iMIS Food modul

Hazards & Fraud





### Control rounds frequency

glass round to be specified for company	monthly	Checking the glass and hard plastic list
cleaning round to be specified for company	daily	Demonstrate proper cleaning in case of production
metal round to be specified for company	daily	At least end of day, before products have left the plant
Visitor check	quarterly	Assurance of the registration process
CCP OPRP competence questions	quarterly	Assurance of HACCP compliance
control specifications	quarter	Assurance of raw materials, end products, packaging
haccp team meeting	quarterly	Possibly with QAssurance per zoom
internal audit	annual	By company itself, in addition to other internal audits, for verificati
monthly round	monthly	Assurance of the functioning of the management system

#### Case

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There are several possible threats and points of attention named in the prerequisite program. The applicable aspects are controlled by procedures, instructions, and the associated measures. weekly

Because of the fact that they originate from the prerequisite program or that the prerequisite program gives guidance for the drafting of these measures, the measures are not guided by the decision tree.

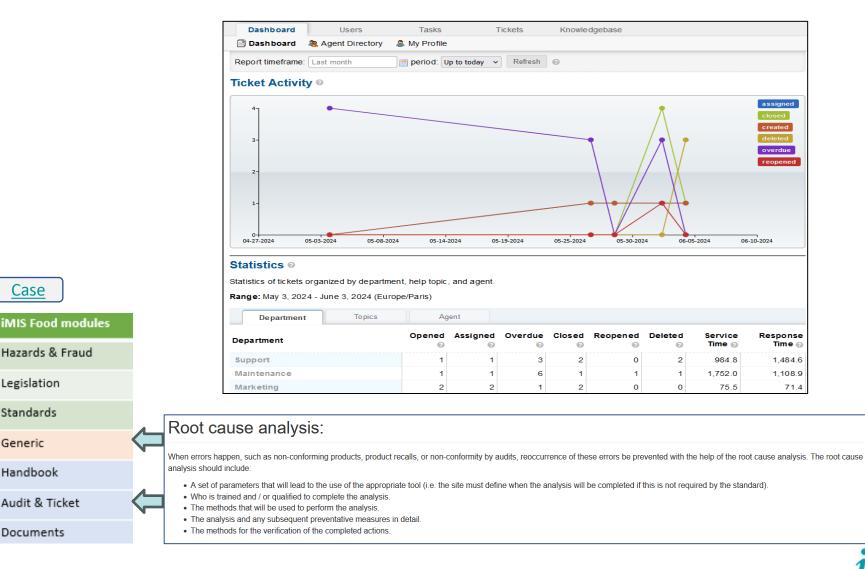
The following measures are in effect within our company and are assessed at least annually or in the case of a new risk or incident in relation to food safety or food defense

Risk from Prerequisite Program	Control by	Standard	Action by abnormalities	Procedure/Instruction
Cross-contamination of product with pathogens with a risk of disease.	Preventing cross-contamination by following company hygiene	Everyone must comply with the applicable regulations	Repeat instruction to personnel, speak to staff. Quarantine extreme products.	Procedure: hygiene Instruction: hygiene control
Cross-contamination with pathogens or hazardous substances due to poor company hygiene in the broadest sense of the word with a risk of disease.	Preventing cross-contamination by following company hygiene	Everyone must comply with the applicable regulations	Repeat instruction to personnel, speak to staff. Quarantine extreme products.	Procedure: company hygiene Instruction: hygiene control
Risk of contamination due to wrong design, wrong materials: not easy to clean, not easy to maintain, FCM not suitable, does not meet equipment requirements. No proper drainage etc.	Furnishing requirements in accordance with BVP. Control rounds, Validation.	Facilities in accordance with requirements.	Modification, maintenance.	Procedures: maintenance, validati BVP check round. Validation repor changes.
Pest feces, bite marks, and etc. Pests can be a carrier of pathogenic agents with a risk of disease.	Good pest control, entry check, and weekly checks on the accumulation of dirt/food products	No pests	Apply additional controls. Instruct personnel. Adjust building technology.	Procedures: pest and entry control Instructions: hygiene control, verm control and entry control
Accelerated spoilage due to the product temperature becoming too high during processing	Preventing temperature increases by controlling the ambient temperature.	See instruction	Quarantine products. Destroy products if temperature is far too high. Control ambient temperature. Decrease ambient temperature. Increase throughput speed. Instruct bersonnel.	Procedures: pests





#### iMIS Food Action List



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Logout

#### **iMIS Food Documents**

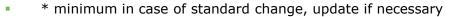
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	01A_System	Options -
	O1B_Quality_handbook	Options -
	O1C_HACCP_study	Options -
	Customers	Options -
ise	O3_Suppliers	Options -
Food modules 32 Folde		Options 👻
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Food modules 32 Folder	Description	
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Food modules ards & Fraud alation alat	Pescription         System folder: The company has a systematic approach to business planning, external visits, and requesting analysis certific:         Quality handbook: HACCP and quality assurance standard related to procedures, instructions, and registration forms are up-         HACCP study: The HACCP study method and result have been recorded (with version number). The HACCP team is described to the study is based on the product specifications, in accordance with which the company wishes to deliver. The risks of third paralso included in the scope of the study.         Customer demands: The company has recorded the most recent customer demands. This concerns terms of delivery, custon specification format. QA relevant information is often also recorded in contracts. Also organize the QA relevant matters with cu	ates. to-date. ed with members and HACCP-related competences. arties (both raw material and outsourced production) ar ner audits, customer questionnaires, and product
Food modules ands & Fraud slation dards Gards A Solution 32 Folder 01-A-System 01-B-Quality- handbook 01-C-HACCP-study	Pescription System folder: The company has a systematic approach to business planning, external visits, and requesting analysis certifice Quality handbook: HACCP and quality assurance standard related to procedures, instructions, and registration forms are up- HACCP study: The HACCP study method and result have been recorded (with version number). The HACCP team is described The study is based on the product specifications, in accordance with which the company wishes to deliver. The risks of third pa also included in the scope of the study. Customer demands: The company has recorded the most recent customer demands. This concerns terms of delivery, custom	ates. to-date. ed with members and HACCP-related competences. arties (both raw material and outsourced production) ar ner audits, customer questionnaires, and product stomer requirements and make them available to the : terms of delivery, supplier audits, supplier
Food modules and Sector 2012 Sector 2012	Pescription System folder: The company has a systematic approach to business planning, external visits, and requesting analysis certifice Quality handbook: HACCP and quality assurance standard related to procedures, instructions, and registration forms are up- HACCP study: The HACCP study method and result have been recorded (with version number). The HACCP team is describe The study is based on the product specifications, in accordance with which the company wishes to deliver. The risks of third pa also included in the scope of the study. Customer demands: The company has recorded the most recent customer demands. This concerns terms of delivery, custom specification format. QA relevant information is often also belong in this folder. Suppliers: The company has recorded the most recent purchasing demands. This concerns general terms of delivery, specific	ates. to-date. ed with members and HACCP-related competences. arties (both raw material and outsourced production) ar ner audits, customer questionnaires, and product stomer requirements and make them available to the : terms of delivery, supplier audits, supplier
Food modules ards & Fraud alation adards cric book	Pescription System folder: The company has a systematic approach to business planning, external visits, and requesting analysis certific: Quality handbook: HACCP and quality assurance standard related to procedures, instructions, and registration forms are up- HACCP study: The HACCP study method and result have been recorded (with version number). The HACCP team is describe The study is based on the product specifications, in accordance with which the company wishes to deliver. The risks of third pa also included in the scope of the study. Customer demands: The company has recorded the most recent customer demands. This concerns terms of delivery, custon specification format. QA relevant information is often also recorded in contracts. Also organize the QA relevant matters with cu organization. Customer satisfaction survey results also belong in this folder. Suppliers: The company has recorded the most recent purchasing demands. This concerns general terms of delivery, specific questionnaires, and product specification format. The newest certifications are known to suppliers. QA relevant information is often	ates. to-date. ed with members and HACCP-related competences. arties (both raw material and outsourced production) ar ner audits, customer questionnaires, and product stomer requirements and make them available to the ; terms of delivery, supplier audits, supplier often also recorded in contracts.

iMIS Documents



### GFSI Reports: all modules needed

- The reports below are delivered during the year in consultation with QAssurance:
- 2.1 Internal audit
- 2.11 HACCP verification
- 2.12 Food Defense analysis TACCP\*
- 2.12 Food Fraud analysis VACCP\*
- 2.13 Emergency plan\*
- 2.14 System Validation
- 2.15 Hygiene photo session
- 2.16 Supplier Assessment
- 2.17 Microbiological scheme
- 2.20 Verification Report
- 2.30 Management review
- 2.40 Recall test
- 2.50 Tracking test
- 2.60 Cross table according to the standard
- 2.70 Audit preparation
- 2.80 Audit guidance
- 2.90 Corrective action handling





#### Case

iMIS Food modules

Hazards & Fraud

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Audit & Ticket

Documents



# GFSI audit: all modules needed

MIS handbook				Search	۹ ۹	• =•	L
en ∄-charts ₽-cross-tables	ISO-TS 22	2.(	002-1				
æ–e-learning æ–general	Titel	Nr	Tekst	beheersing	verwijzi	zing	
⊞-haccp-study ⊞-imis-checklist	Construction and layout of buildings	4					
instructions     instructions-specific     forcedures     procedures-specific     registrations     risk-analysis     scorecards     secifications     reschortations-and-responsi	J	4.1	Buildings shall be designed, constructed and maintained in a manner appropriate to the nature of the processing operations to be carried out, the food safety hazards associated with those operations and the potential sources of contamination from the plant environs. Buildings shall be of durable construction which presents no hazard to the product. NOTE For example, roofs should be self-draining and not leak.	s gecontroleerd via de iMIS iMIS Audit controlerondes.		udit controlero	ron
9) tasks-authorizations-and-responsi - 32-folder - broken-links - exportpdf - index - standards	Environment	4.2	2 Consideration shall be given to potential sources of contamination from the local environment. NOTE Food production should not be carried out in areas where potentially harmful substances could enter the product. The effectiveness of measures taken to protect against potential contaminants shall be periodically reviewed.	De omgeving heeft geen negatieve invloed op de veiligheid en kwalite Jaarlijks wordt dit meegenomen in de management review en verificat	0	jement review atie	N
P DUTCH Translated from English	Locations of establishments	4.3	<sup>3</sup> The site boundaries shall be clearly identified. Access to the site shall be controlled. The site shall be maintained in good order. Vegetation shall be tended or removed. Roads, yards and parking areas shall be drained to prevent standing water and shall be maintained.	De terreingrenzen zijn duidelijk aangegeven. De gebouwen en het terrein zijn goed onderhouden. Water loopt voldoende snel weg via het riool. De vegetatie wordt kort gehouden. Dit alles wordt gecontroleerd met de iMIS Audit controlerondes.	iMIS Au	udit controlero	roi
=-cross-cables	Layout of premises and workspace	5					
	General requirements	5.1	Internal layouts shall be designed, constructed and maintained to facilitate good hygiene and manufacturing practices. The movement patterns of materials, products and people, and the layout of	De routing is logisch van opzet. Door de gehanteerde route worden kruisbesmettingen voorkomen, zie ook de plattegronden. In de route is aandacht voor personeel, product, machines,	Plattegro iMIS qui		

Legislation

Case

iMIS Food modules

Hazards & Fraud

Standards

Generic

Handbook

Audit & Ticket

Documents





# iMIS Food Helpdesk the support for (un)announced audits

- Content
  - ✓ Legislation & HACCP study
  - ✓ Quality standards
  - Inspection rounds
  - Labelling & Specifications
  - Available 24 hours a day for recalls

#### Case

#### iMIS Food modules

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#### Software

- ✓ Support in using the software
- Remote takeover of the PC
- Import and export of data
- Company-specific reports
- Login, users, passwords







# Server: calendar, webforms and documents

			< February	2024 >						M
	Handbook		Saturday 21	7 Sunday 21	Monday 2	Tuesday	Wednesday 30 31	Thursday	Friday 5:17 System Update 0% 8:5:28 disk and hardplastic 92%	Legend: • futureh • futureh Printer Fin Preference Jump to
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	Documents		• 10:02 gc daily operation check 82% Site	10:04 gc daily operation check 89%	1 10:08 qc daily operation check 55% 5ite 10:28 qc daily operation check 55% fite 0 gc daily operation check 56% Rotterdam	<ul> <li>10:09 qc daily operation check 90 Rotterdam</li> <li>10:12 qc daily operation check 92 Site</li> </ul>	13 • 10:13 qc daily operation check 03% Site %	10:15 qc daily operation check 74%	5 10	Go
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6						1.11				30 31
	Calendar	IMIS Webforms							Search	۹ ۹
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	iMIS Audit Action List		Company Name Goal				t is forbidden for una mandatory for visitor read and sign them b registration form (visi 1. Visitors who have been l pose a danger to food s	authorized persons s to follow the com before entering the itor list) to register in contact with, or suffer fro afety must NOT enter prod	s to enter the product apany rules. To do the production. This is a visit.	nis, they mus done on the er illnesses that m
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Case

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