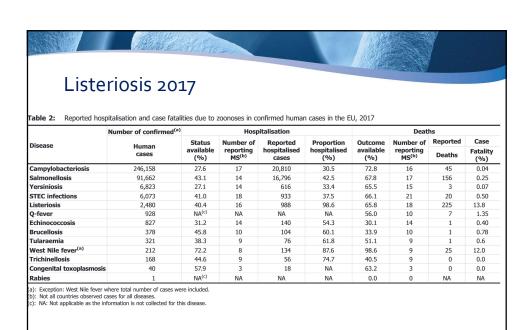


PROF. JOHN HOLAH, TECHNICAL DIRECTOR

Fagdag om Listeriakontroll,13th November 2019, Gardermoen, Norway



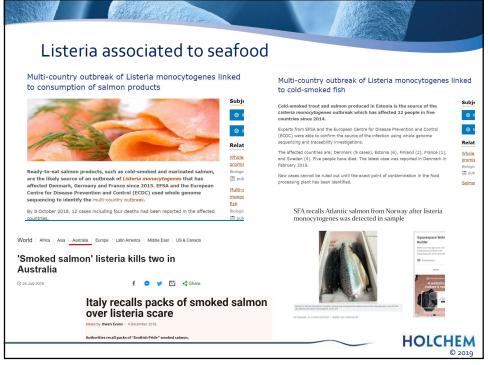
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The European Union summary report on trends and sources of zoonoses, zoonotic agents and food-borne outbreaks in 2017 EFSA Journal 2018;16(12):5500



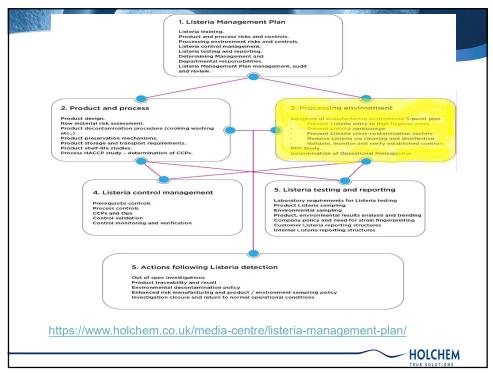
_	9	ops a	ected			
	2017	2016	2015	2014	2013	Data source
RTE food RTE food – occurrence	(%) by deta	action metho	d (number of	f tostod same	ales by detec	tion method:
number of reporting M		ection metho	a (namber o	testeu sam	nes by detec	don metriou,
Fish and fishery products	6.0% (n = 6,730; 22 MS)	5.1% (n = 2,918; 22 MS)	3.2% (n = 4,658; 22 MS)	5.8% (n = 3,436; 16 MS)	5.1% (n = 3,479; 20 MS)	EFSA
Meat and meat products (beef, pork, broiler and turkey meat)	1.8% (n = 22,544; 19 MS)	3.3% (n = 15,161; 23 MS)	2.8% (n = 16,789; 21 MS)	2.1% (n = 67,215; 18 MS)	3.4% (n = 44,977; 21 MS)	EFSA
Soft and semi-soft cheeses made from raw or low-heat-treated milk	0.9% (n = 6,117; 17 MS)	2.6% (n = 853; 15 MS)	1.4% (n = 730; 13 MS)	1.0% (n = 2,573; 13 MS)	4.2% (n = 2,542; 13 MS)	EFSA
Hard cheeses made from raw or low-heat-treated milk	0.1% (n = 5,039; 15 MS)	1.0% (n = 509; 9 MS)	1.3% (n = 858; 11 MS)	0.2% (n = 10,175; 9 MS)	0.7% (n = 1,609; 12 MS)	EFSA
Fruit and vegetables	0.6% (n = 1,773; 17 MS)	0.7% (n = 1,043; 16 MS)	2.1% (n = 1,456; 17 MS)	3.0% (n = 1,503; 17 MS)	2.1% (n = 1,991; 15 MS)	EFSA
Salads	4.2% (n = 902; 14 MS)	1.9% (n = 1,042; 14 MS)	1.9% (n = 1,238; 13 MS)	1.1% (n = 1,154; 15 MS)	2.4% (n = 1,822; 14 MS)	EFSA

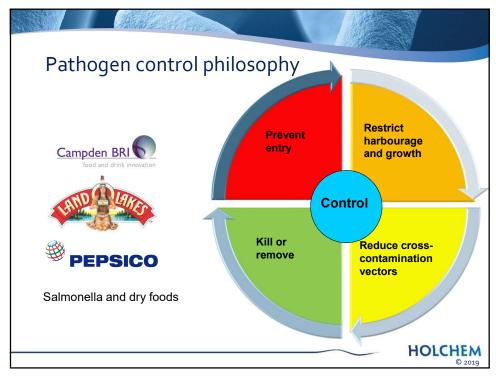


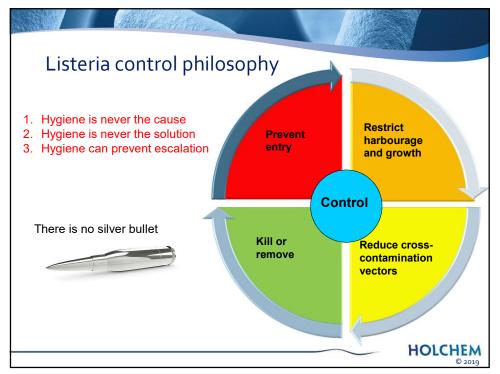
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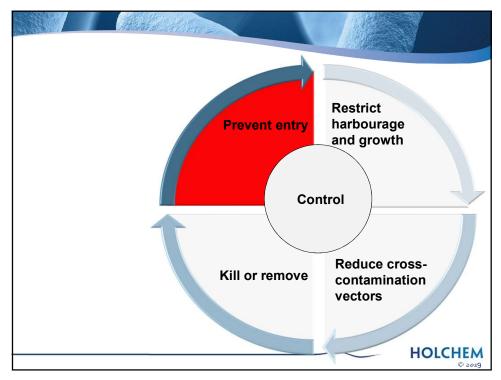


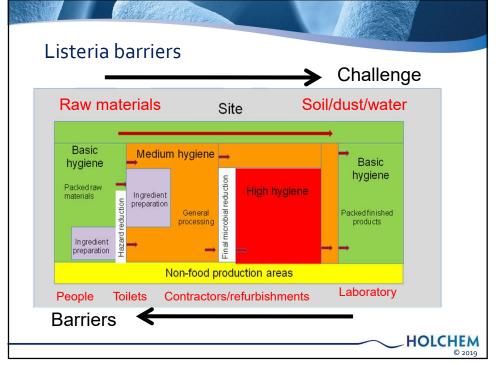
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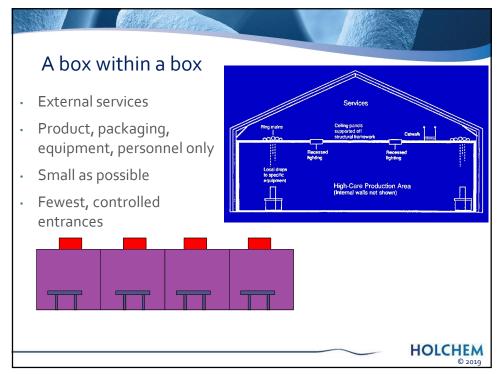


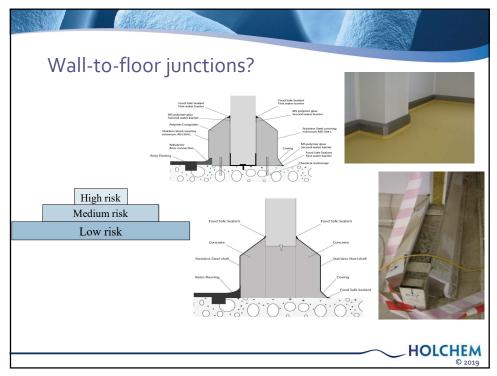


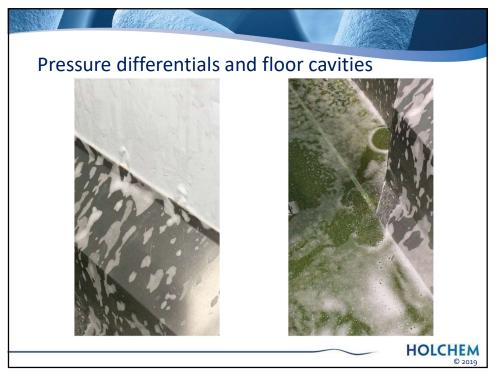


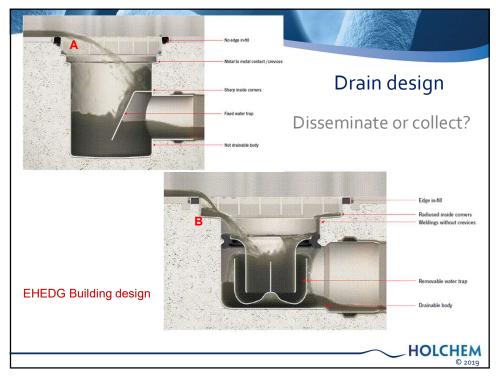


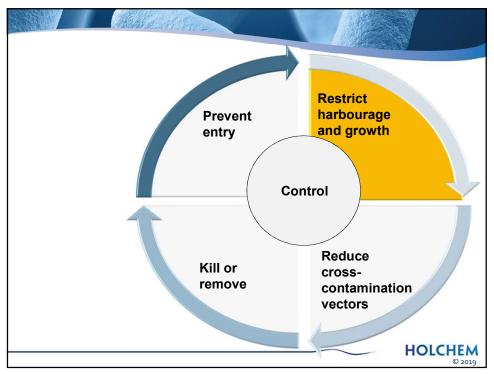




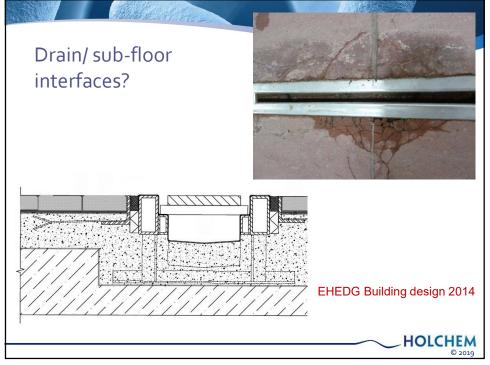


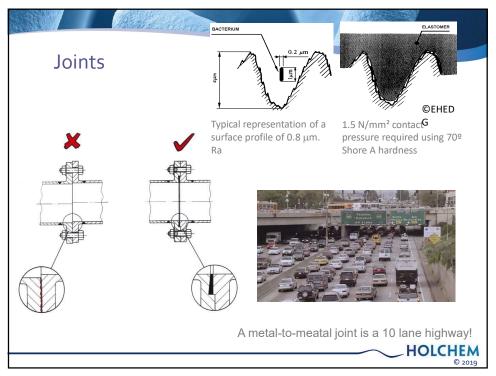


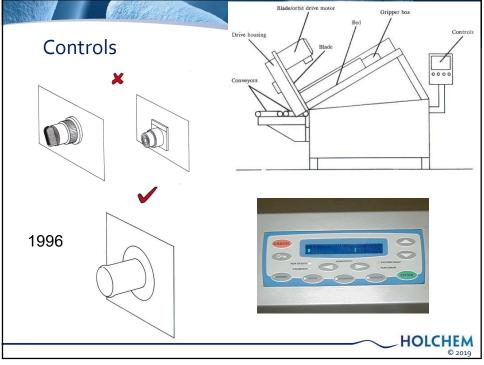




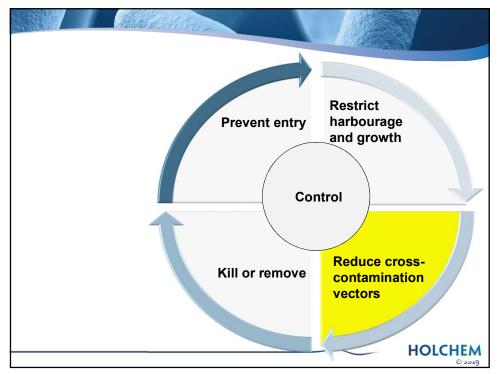


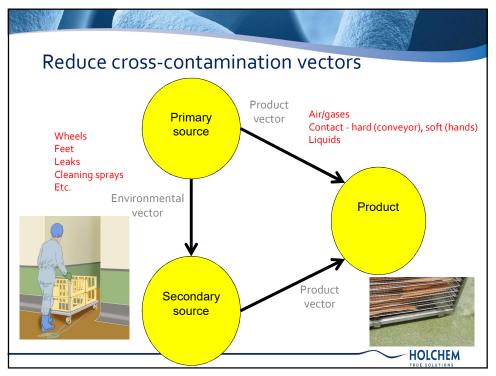


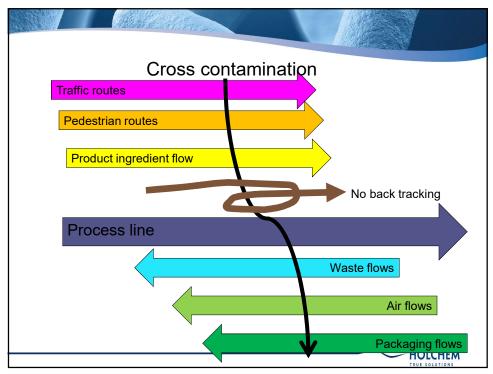


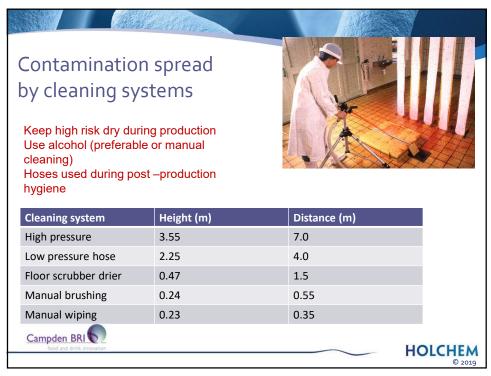


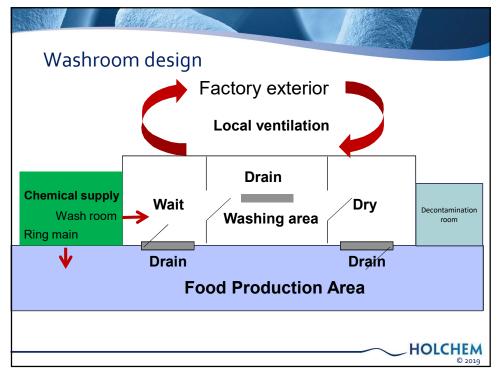


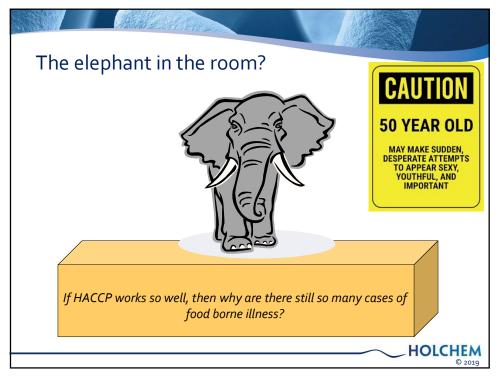












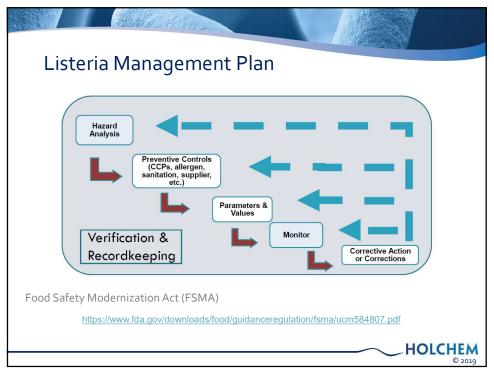


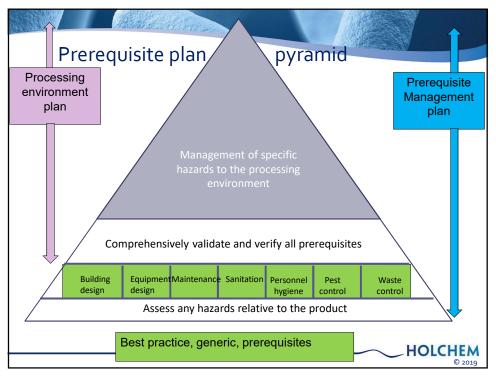
Proposed hazard analysis model

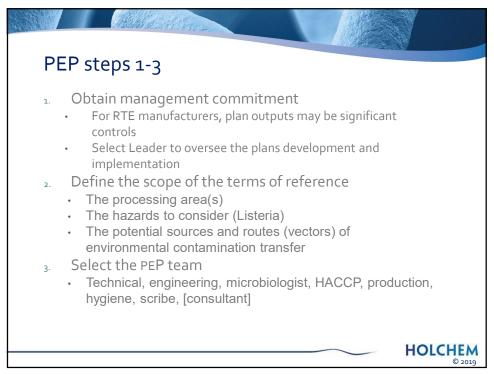
- A cheese and onion sandwich is about to be manufactured after a peanut butter sandwich. A between-batch clean has been undertaken
 - An operative on the line is not wearing a hair net.
 - The line still has peanut residues on it
 - · What would you do! Would you stop the line?
- As part of Prerequisite Management, develop a Processing Environment Plan
- Reduce terminology use the same 14 principles of the HACCP plan (Campden BRI Guideline No.42)
- US approach is HARPC (Hazard Analysis and Risk-Based Preventive Controls)
- https://www.foodsafetymagazine.com/enewsletter/haccp-vsharpc-a-comparison/



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PEP steps 4-5

- Describe the environment
 - Size, zones, boundaries, barriers, services, air flows, temperatures, humidity, people flows, transport flows, liquid and solid waste flows
 - Production periods, sanitation procedures, shut downs
 - Historical sampling and observational data e.g. microorganisms, pests, glass
- 5. Identify intended use
 - Further treatments: removal, reduction or growth
 - Consumer: target consumer group



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PEP steps 6-7

- 6. Construct flow diagram
 - Base layer of processing area, equipment and services
 - Overlaying plans of e.g. air flows, people movement, transport movement, waste flows
- 7. On-site confirmation of flow diagram
 - Sign off diagram(s) as an accurate representation of the processing area and activities

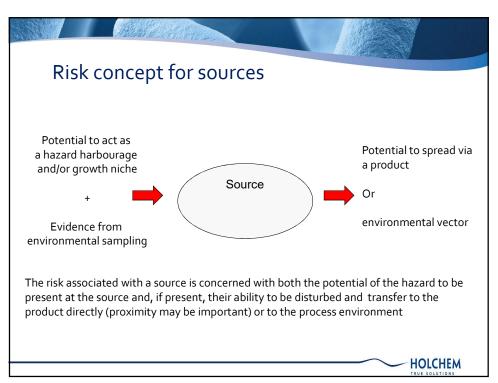


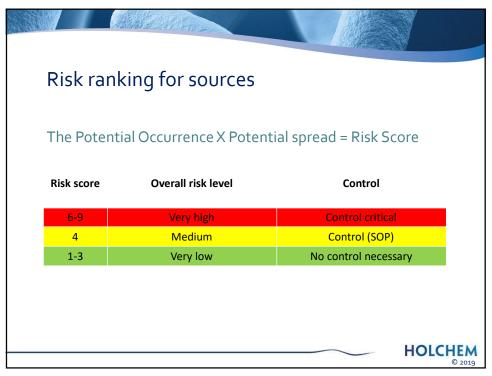
PEP step 8

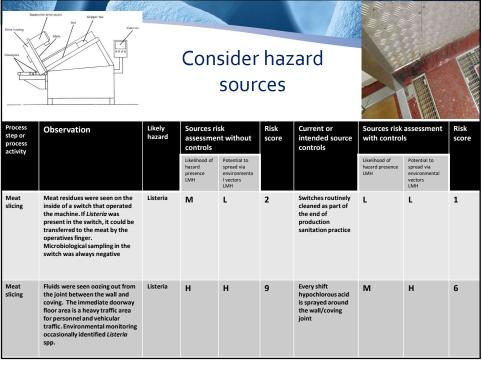
- 8. List all potential hazards, conduct a hazard analysis and consider any measures to control the identified hazards
 - Undertake factory study by LMP team to investigate the defined sources and vectors of environmental and product contamination
 - Sources:- history internal, history external, extensive microbiological sampling.
 - Vectors:- visual assessments and discussions with staff re: all line interventions; physical measurements e.g. of airflow
 - Determine (any) current control options for the identified sources and vectors. Suggest additional/new control options.
 - Assess the need for control validation and verification

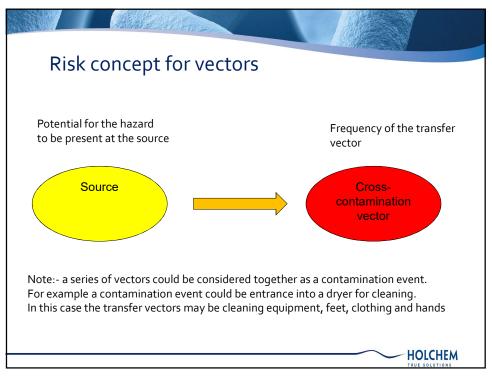


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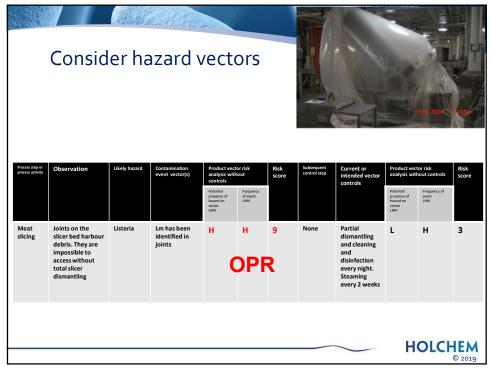




- 9. Determine Operational prerequisites
 - Determine whether any hazards will be removed by subsequent control steps
 - Determine whether, in the absence of controls, any risks are so significant that they could be described as Operational Prerequisite
 - An OP should have the same 'status' as a CCP



Consider hazard sources									
Process step or process	Observation	Likely hazard	Sources risk assessment without controls		Risk score	Current or intended source controls	Sources risk assessment with controls		Risk score
activity			Likelihood of hazard presence LMH	Potential to spread via environmenta I vectors LMH			Likelihood of hazard presence LMH	Potential to spread via environmental vectors LMH	
Meat slicing	Meat residues were seen on the inside of a switch that operated the machine. If <i>Listeria</i> was present in the switch, it could be transferred to the meat by the operatives finger. Microbiological sampling in the switch was always negative	Listeria	М	L	2	Switches routinely cleaned as part of the end of production sanitation practice	F	PR	1
Meat slicing	Fluids were seen ozing out from the joint between the wall and coving. The immediate doorway floor area is a heavy traffic area for personnel and vehicular traffic. Environmental monitoring occasionally identified <i>Listeria</i> spp.	Listeria	н	Н	9	Every shift hypochlorous acid is sprayed around the wall/coving joint	CA	" \PEX	6
								HOLCH	



PEP steps 10-11 10. Establish control or operating limits

- Likely to be factory based, potentially critical (but unlikely)
 - Measurable or observable relative to control
- 11. Establish a monitoring system
 - Detection of any loss of control prior to food being cross contaminated
 - E.g. ATP, protein, allergen, pH, chemical concentration, temperature
 - Establish any national reference methods
 - Authority, competence, training



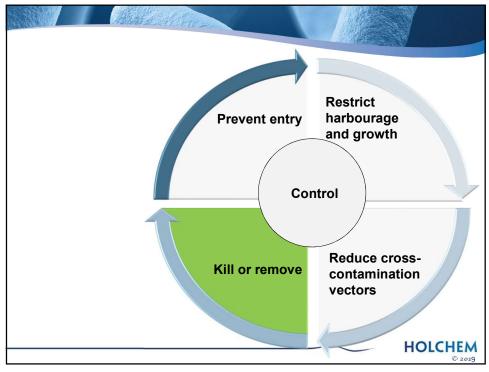
PEP step 12-14

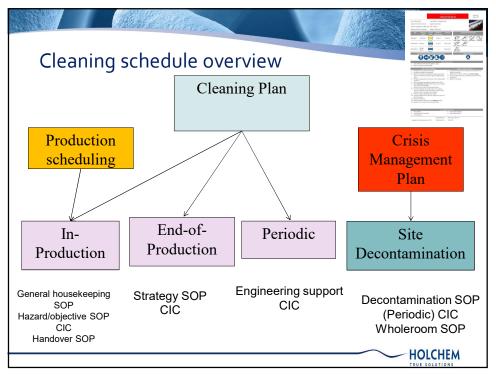
- 12. Establish a corrective action plan
 - Defined actions, product hold procedures
- 13. Verification
 - Validation of plan
 - Validation of any essential controls
 - Environmental sampling plan
 - · Verification by e.g. microbiological testing
 - Audit, trend analysis, customer complaints
 - Review any process or processing environment changes
- 14. Establish documentation and record keeping
 - All records to demonstrate effective control to customers



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OPR management sheet						11 200C 1 COM		
Process step or area	Likely hazards	Source or Contamination event vector(s)	Control measure(s)	Operating limit(s)	Control monitor(s)	Control verification(s)	Corrective action(s)	Records
M eat slicing	Listeria	Transfer of Lm from slicer bed to product during slicing	Cleaning until visual cleanliness ATP testing prior to disinfection Attachment of thermocouples Encasement of slicer and ancillary components in plastic bag Overpressure of sensitive components Steaming until thermocouples read 70°C and then for a further 30 minutes Drying until moisture free	ATP <150 RLU 70°C 30 min	Visual assessment of freedom of debris ATP assessment prior to disinfection Monitoring of time and temperature	Microbiological verification of slicer bed Thermocouple calibration Timer calibration Steam pressure	Staff retraining Recalibration	Visual cleanliness, ATP and microbiological records Temperature and time records Steam pressure records Calibration records
								© 2019





End of production

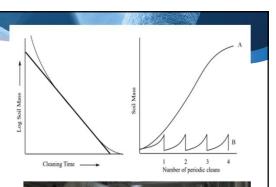
- · Handover from production to hygiene
 - · Defined cleaning window
 - · Removal of all product and packaging
- Sequence
 - Gross solids removal, rinse, Listeria sources (e.g. drains), environment, equipment
- · Chemical rotation
 - · pH not disinfectants
- Cleaning validation worst case scenarios
 - Hygienic design dead zones, most difficult to remove soil, minimal acceptable cleaning programme



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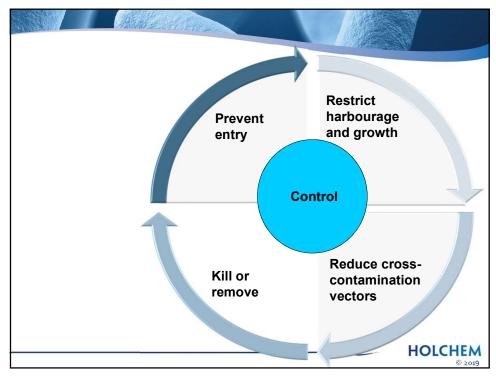
Periodic cleaning

- Cleaning is not 100% efficient so soil will accumulate with time
- Periodic cleans are required
 - Additional energy
 - Additional strip down
 - Alternative chemicals e.g. scale removal (acid clean)
 - Heat





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Sampling principles

All samples must have a purpose

• What is your corrective action if the sample is positive?

Size matters

- Large sponge/wipe samples to maximise detection e.g. on collectors
- Small swabs to focus on identified areas that are difficult to clean

Validated sampling techniques

- Listeria method ISO 17025 accredited
- · Laboratory undertakes proficiency tests
- · Ability to neutralise disinfectants
- · Ability to maintain captured pathogens viability





• Sample around barriers during the day e.g. personnel changing, product entry tunnels, air supply and verify performance

Is everything under control during production?

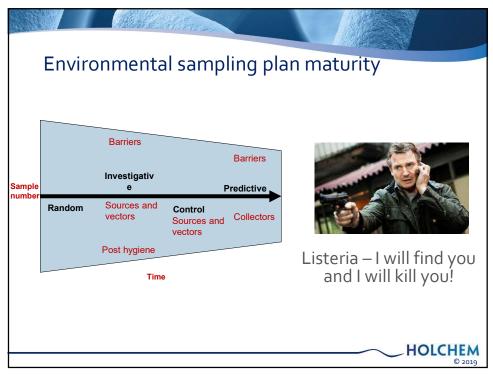
- Sample known/suspected sources and transfer vectors are they in control?
- · Sample collector points e.g. footwear, tote wheels, cleaning equipment, drains – is there any evidence of the presence of a pathogen?
- · Investigative studies if yes

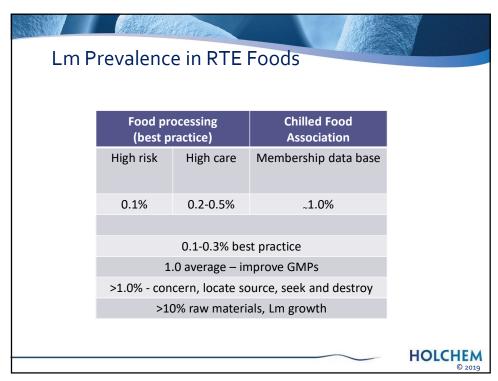
Do we have pathogen free processing equipment for subsequent production?

· Verify cleaning and disinfection performance



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Sampling site vs Lm detection rate Sample site Sample position Sample position Barriers 10 Source/vector control 10 Collectors 10 Cleaning verification 70 Detection rate <0.1%

Sampling site vs Lm detection rate

Sample site	Sample position	Sample position
Barriers	10	10
Source/vector control	10	10
Collectors	10	70
Cleaning verification	70	10
Detection rate	<0.1%	>0.2%

If Lm is from an environmental source env. Lm 2-3x product Lm If Lm is from a raw material source product Lm> env Lm



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