## Level 4 Award in Managing the HACCP System for the Meat Industry Marking Guide, Specimen Paper B



1a)	Explain why the process flow diagram should be verified.	To ensure that To ensure that happens in rea To ensure that captured/unde Doing so makes properly consider	all situations/situations rstood s sure that all food safet dered and evaluated. o could result in hazards	is correct ately reflects what are cy hazards can be	5 Marks
1b)	Describe how this should be done for the venison HACCP plan.	Possibly large variability in conditions at time of killing Therefore PFD verification will require visits into the field/estate, under a range of conditions (weather, seasons, locations, different huntsmen) Peak demand periods need to be considered Observe whole of the operation Identify food/people flows Consider drainage/air flows Ask someone off the HACCP team to do it Sign and date the PFD as verified, amending if necessary Record, not the range of conditions observed for use in the hazards analysis		5 Marks	
2a)	Referring to the LABELLING process step and the information about Clostridium botulinum:	PROCESS STEP	Food Safety Hazards and Cause	Control Measure	5 Marks
	Identify a significant food safety hazard and its cause together with suitable control measure(s).	Labelling of - vacuum packed meat	Growth of Clostridium botulinum (and production of heat	Correct application of the date code according to the	

b)	Decide whether the process step is critical to safety. Use the supplied CCP decision tree to justify your answer.	Question 2 – NO to eliminate the Question 3 – YE unacceptable le Question 4 – NO	e hazard S, the identified hazard evels during the shelf li D, no further process st e the toxin is heat stab	ot specifically designed d could increase to fe tep will eliminate the	5 marks
3a)	Referring to your answer above, suggest an appropriate:  Monitoring plan	PROCESS STEP Labelling of - vacuum packed meat	Monitoring Plan  WHO – Trained oper WHEN – at start and hourly in between HOW– Take a label f machine. Check that displays the critical I days). Check that the instructions state "st RECORD– stick the p CCP monitoring shee and sign	rom the date coding the label clearly imit shelf life (+8 e consumer storage tore below 8C"	5 Marks
b)	Corrective action plan	PROCESS STEP Labelling of - vacuum packed meat	report to Supervisor IMMEDIATE: Product product from the line check) must be held INTERMEDIATE: Engi investigate and fix da	/trained operative to immediately tion line halted, all e (from the last good in quarantine.	5 Marks

		line re-started with increased monitoring frequency. Each date label for food in quarantine to be visually checked. Any with clear date can be released, any with unclear date label must have date label removed and be passed back through the date labeller  RECORD: Full record of all actions taken to be made  PREVENTION: Conduct root cause analysis.  Why did printer fail? PPM adequate etc.	
4a)	To improve competitiveness in the export market, the HACCP team want to know if they can safely increase the shelf life of the vacuum-packed product.  What factors should they consider?	Scientific literature, Industry guidance, Government guidance/advice (e.g. FSA, ACMSF). If storage instructions (refrigeration) could be changed to 3C or below, if they could change the product/product formulation, what shelf life their competitors are giving, challenge testing, micro-modelling, shelf life trials	5 Marks
b)	The team decide to extend the product shelf life by two days. How should this change be implemented?	The team decide to extend the product shelf life by two days. How should this change be implemented?	5 Marks
5a)	The OV conducts an audit to verify compliance with Official Controls and asks to see all documents and records relating to the labelling process step. She is shown some recent monitoring records and discovers that the old document is still being used (specifying the 8 day product shelf life). The OV is concerned that document control is being poorly managed.  Apart from the monitoring document,	Standard Operating Procedure – do they make sense? Are operators are following them? Actual product labels – are they are clearly printed and have the correct date? Maintenance records for the date coding machine – to show that the machine will produce clear labels and is less likely to fail Training records for production staff – to demonstrate competence and understanding in the use of the date coding machine On the job "sign off sheets" - showing staff competency in use of date coding machine	5 Marks
	identify 5 other records or documents the OV may request and for each one explain how it can be used to assess whether the process is under control.	HACCP chart for the process – to check if HACCP principles have been followed (e.g. correct identification of hazards, correct CCP decision etc?)	

		Validation evidence – e.g. results of challenge testing. Will the food safety hazards be effectively controlled?  Terms of reference – showing details of industry/technical guidance referred to  Corrective action records – are they being taken? Are lessons learned?  Internal auditing records/reports – have CAR's been closed out? Are there repeated issues?	
5b)	Explain why good document control is required for an effective food safety management system.	Document control is a formal system for creating, modifying, and reviewing, issuing, distributing and ensuring accessibility of documents.  Makes sure the food safety management system is "kept live". Ensures that all documents are trusted, up to date and reliable.  Documents are checked before issue and formal approval means they carry the correct level of authority.  Reduces the risk of errors and therefore reduces food safety risk	5 Marks