

Code of Practice: Processing of Bee Products Part 5: RMP Templates

RMP Template for the Processing of Honey and Dried Pollen

5 RMP Template for the Processing of Honey and Dried Pollen

This RMP template applies to businesses that are involved in the processing of honey or dried pollen.

The *Guidelines for Completing the RMP Template* should be referred to when completing this template.

The RMP template starts on the next page. This page is not part of the RMP.

1.	Business Identific	ation	
Rus	siness ID:		RMP No.:
		using an Address and Contact Pater	
2.	Operator Name, B	usiness Address and Contact Deta	alls
Ful	l legal name (Company,	sole trader, partnership):	
Tra	ading name (if different):		
Phy	ysical address(es) of pre	emises:	Phone No:
			Fax No:
			E maile
			E-mail:
Pos	stal address (for commu	nication):	
1 00	otal address (for comma	modion).	[] I give consent to being provided electronic information.
3.	Responsible Perso	on	
Rol	le	Name, position or designation	Contact Details (if different from above)
Day-to-day Manager of the RMP			
4.	Scope of the RMP		
[1 The physical hour	ndaries of the RMP are shown on the atta	tached site plan
	1 The physical book	idented of the riving are shown on the atte	acred site plan.
The	e RMP covers the follow	ing processes or activities:	
[] Extraction of hon	ey	[] Melting and moulding of bees wax
[] Processing and p	packing of liquid or creamed honey	[] Drying, cleaning and packing of dried pollen
[] Cutting and pack	ing of comb honey	
]] Storage of honey		[] Other (specify)
	e following products or der a different RMP or u		boundaries of the RMP are excluded because they are covered
	Product or activity	: Covered under:	
		[] Another RMP N	No [] Food Act
		[] Another RMP N	No [] Food Act

5. Product Description			
Products	Bulk honey	Liquid or creamed honey	Comb honey
Intended consumer	Humans (general public)	Humans (general public)	Humans (general public)
Intended use of product that leaves RMP	 Further processing and packing to liquid/creamed honey or other honey products Ingredient for preparation of other foods 	Ready-to-eat Ingredient for preparation of other foods	 Ready-to-eat Ingredient for preparation of other foods
Regulatory limits	None	None	None
Other regulatory requirements specific to product	Food Standards Code 2.8.2 - • Reducing sugars ≥ 60% • Moisture ≤ 21%	Food Standards Code 2.8.2 - • Reducing sugars ≥ 60% • Moisture ≤ 21%	Food Standards Code 2.8.2 - • Reducing sugars ≥ 60% • Moisture ≤ 21%
	AP (Residue Specification) Notice 2004 - Specified chemical substances in honey ≤ maximum permissible levels	AP (Residue Specification) Notice 2004 - Specified chemical substances in honey ≤ maximum permissible levels	AP (Residue Specification) Notice 2004 - Specified chemical substances in honey ≤ maximum permissible levels
	Every consignment of honey must be provided with an <i>Apiarist and Beekeeper Statement</i> (i.e. Harvest declaration) and comply with HC Spec 108.	Every consignment of honey must be provided with an <i>Apiarist and Beekeeper Statement</i> (i.e. Harvest declaration) and comply with HC Spec 108.	Every consignment of honey must be provided with an <i>Apiarist and Beekeeper Statement</i> (i.e. Harvest declaration) and comply with HC Spec 108.
Labelling	Labelling of transportation outers as specified in HC Spec 32.	Labelling of retail packs as specified in the Food Standards Code. Labelling of transportation outers as specified in HC Spec 32.	Labelling of retail packs as specified in the Food Standards Code. Labelling of transportation outers as specified in HC Spec 32.

	5. Product Description (continued)					
eeswax	Spilt honey, downgraded honey (e.g. fermented)	Dried pollen	Others			
umans (general public)	Animals	Humans (general public)				
Further processing into products for pharmaceutical use and manufacture of cosmetics Further processing into comb foundation	Feed for bees and other animals (e.g. horses).	Ready-to-eat Ingredient for preparation of other foods & dietary supplements				
one	None	None				
/A	N/A	Every consignment of pollen must be provided with an <i>Apiarist and Beekeeper Statement</i> and comply with HC Spec 108.				
abelling of transportation outers s specified in HC Spec 32.	Labelled "Not for Human Consumption"	 Labelling of retail packs as specified in the Food Standards Code including an advisory statement as required by Standard 1.2.3. Labelling of transportation outers as specified in HC Spec 32 				
or /A	Further processing into products for pharmaceutical use and manufacture of cosmetics Further processing into comb foundation ne	Further processing into products for pharmaceutical use and manufacture of cosmetics Further processing into comb foundation None None N/A Labelled "Not for Human	Purther processing into products for pharmaceutical use and manufacture of cosmetics Further processing into comb foundation None None			

Risk Management Programme

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6.	6. Process Description							
Bulk honey		Liquid or creamed honey	Comb honey	Beeswax				
[] Receiving supers	[] Receiving of bulk honey	[] Receiving of honey supers	[] Collection of cappings and other wax material				
[] Holding in hot room/ store room	[] Storage	[] Storage of supers	[] Separation of honey from cappings				
[] Deboxing	[] Cleaning drum external surface	[] Deboxing	[] Melting of wax				
[] Uncapping	[] Heating in hot room	[] Removal of wires	[] Filling of wax into moulds				
[] Pricking/loosening	[] Pouring honey into vats/tanks	[] Inspection of combs	[] Cooling				
[] Extraction	[] Heating using heat exchanger	[] Cutting of combs	[] Dispatch				
[] Transfer through sump	[] Filtering	[] Packing & labelling					
[] Heating using heat exchanger	[] Creaming	[] Freezing					
[] Spinning	[] Holding in tanks	[] Dispatch					
[] Pumping into tanks & straining	[] Packing and labelling						
[] Holding in tanks	[] Storage						
[] Filling of honey into drums	[] Dispatch						
[] Labelling/marking of drums							
[] Storage							
[] Dispatch							

Date:

Date:

7. External Verification

Verifier's Freedom and Access to carry out Verification Functions (RMP Specifications 2003, clause 15)

I authorise my contracted verifier to have the freedom and access necessary to allow him/her to carry out verification functions and activities, including —

- (a) having access to all parts of the premises or place and facilities within the physical boundaries of, or relating to, the risk management programme; and
- (b) having access to all documentation, records and information relating to, or comprising, the risk management programme (including records held in electronic or other form); and
- (c) having freedom to examine all things necessary and open any containers, packages and other associated things to inspect their contents; and
- (d) having freedom to identify or mark any animal material, animal product, equipment, package, container or other associated thing; and
- (e) having freedom to—
 - (i) examine and take samples of any animal material, animal product or any other input, substance, or associated thing which has been, is, or may be in contact with, or in the vicinity of, any animal material or animal product; and
 - (ii) test, or analyse, or arrange for the testing or analysis of such samples; and
 - (iii) order retention of materials including animal material, ingredients, animal product, packaging or equipment pending testing results and decisions on disposition; and
- (f) having authority to detain any animal material and animal product or other relevant things in the event of non-compliance with the risk management programme where there may be significant risk to fitness for intended purpose of animal product or suitability for processing of animal material; and
- (g) having authority to intervene and direct a temporary interruption of processing in cases of significant risk to fitness for intended purpose of animal product or suitability of animal material for processing until the cause of the risk has been remedied.
- [] A letter has been received from the verification agency confirming they will verify the RMP at all sites covered by this RMP.

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Document	Documents from the COP		Operator's own documents based on the COP		Operator's own documents for additional products/processes/ procedures		Person responsible for Implementation
	Reference	Date	Reference	Date	Reference	Date	
Main part of RMP (this document)	N/A		Completed RMP template				
GMP Supporting Systems:	***************************************						
Design and construction of buildings, facilities and equipment							
Potable water							
Cleaning and sanitation							
Personnel competency, health and hygiene							
Control of chemicals							
Pest control							
Packaging materials (specifications, handling and storage)							
Receipt and processing of honey and dried pollen							
Document control and record keeping (including inventory control)							
Recall of products							
Operator verification							

Document	Documents from the COP		Operator's own documents based on the COP		Operator's own documents for additional products/processes/ procedures		Person responsible for Implementation
	Reference	Date	Reference	Date	Reference	Date	•
HACCP Application							
Identification of risk factors related to wholesomeness and labelling							
Other documents:							
Site plan of physical boundaries Letter from Verification Agency							
Assessment of Water Supply Status (only necessary for							
Agency Assessment of Water Supply Status (only necessary for own supply)							

9.	Cor	firmation				
[]	I confirm that all of the documents listed in Section 8 are appropriate for my operation.				
[]	I confirm that all facilities and equipment necessary to implement the RMP are available and ready to operate.				
[]	I confirm that the RMP, including all supporting systems, has been authorised by me.				
[]	I confirm that the RMP will be implemented as written, including all relevant parts of the code of practice.				
1						
Sig	Signature of Operator or Day-to-day Manager of RMP: Date: / /					