FOOD INDUSTRY - PRODUCT INFORMATION FORM

VERSION 5.0 - released 15 May 2012



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mouse right click on box bitmap edit VARRANTY: This document is intended as a guide only: legal requirements are contained in the Food tandards Code and relevant food legislation and other applicable laws. The information in this document hould not be relied upon as legal advice or used as a substitute for legal advice. You should exercise your own kill, care and judgement before relying on this information in any important matter.

1 CONTACT DETAILS & DECLARATION						
SUPPLIER'S	Poppy Seed	SPECIFY COUNTRY				
PRODUCT NAME	Poppy Seed	IMPORTED INTO				
SUPPLIER'S	ın/a	SPECIFY COUNTRY				
PRODUCT CODE	II/a	EXPORTED FROM	Australia			
BARCODE -	n/a	SPECIFY IMPORT				
UNIT GTIN	III/a	TARIFF CODE				

1.1 SUPPLIER INFORMATION

	OUT I LILIX IIVI OKIMATION						
	COMPANY NAME	Sun Pharmaceutical Indu	Sun Pharmaceutical Industries (Australia) Pty Ltd				
	BUSINESS NUMBER (ABN)	64130119603					
TRADING NAME		Sun Pharma Controlled S	Substances Division				
BUSINES ADDRES		14 Henry Street	L	atrobe			
	STATE / COUNTRY / POST CODE	Tasmania	Australia		7307		
POSTAL	POST ADDRESS / SUBURB	PO Box 189	L	atrobe			
ADDRES	CITY / COUNTRY / POST CODE	Tasmania	Australia		7307		
KEY CON	TACT NAME	Tiago Tomaz					
FOR QUE	RIES POSITION TITLE	Head of Crop, Research & Operations					
EMAIL ADDRESS		tiago.tomaz@sunpharma.com					
	PHONE	61 (0)3 6426 5700	FA	AX			
DATE FORM COMPLETED		08-April-2020	ISSUE DA	ТЕ			
	DOCUMENT NO:		ISSUE NUMBE	R			

1.2 MANUFACTURING INFORMATION

Provide details where the manufacturer or site location differ to above:

		COMPANY NAME	Sun Pharm	naceutical Industries (Australia) Pty Lt	d	
SITE:	#1	NUMBER / STREET / SUBURB	14	Henry Street		Latrobe	
		STATE / COUNTRY / POST CODE	Tasmania		Australia		7307
		COMPANY NAME					
SITE:	#2	NUMBER / STREET / SUBURB					
		STATE / COUNTRY / POST CODE					
		COMPANY NAME					
SITE:	#3	NUMBER / STREET / SUBURB					
		STATE / COUNTRY / POST CODE					

If more than three manufacturing sites, provide additional site information in Section 8.2

1.3 CONTACT DETAILS FOR TECHNICAL & ALLERGEN INFORMATION

Please specify the contact details if further information related to technical or allergen information is needed:

	<u> </u>			
NAME	Tiago Tomaz / Cassandra Amos			
JOB TITLE	Head of Crop, Research & Operations / Qualitry Assurance Officer			
EMAIL	tiago.tomaz@sunpharma.com / cassandra.amos@sunpharma.com			
TELEPHONE - WORK	61(0)3 64265700	TELEPHONE - MOBILE	0449 953 137 (Tiago)	

1.4 SUPPLIER DECLARATION AND WARRANTY

The Supplier -

- 1) certifies that this product complies with the Australia New Zealand Food Standards Code; and, in addition to the information provided specifically in this form, and without limitation to compliance with any other part of the Code, that the product complies with:
 - (a) Standard 1.3.4 Identity and Purity
 - (b) Standard 1.4.1 Contaminants & Natural Toxicants
 - (c) Standard 1.4.2 Maximum Residue Limits in Food (In Australia), or
 - (d) Maximum Residue Limits of Agricultural Compounds, Mandatory Food Standard 1999 (and subsequent amendments) issued under sections 11C and 11Z of the Food Act 1981 in New Zealand
 - (e) Standard 1.4.3 Articles & Materials in Contact with Food
 - (f) Standard 1.4.4 Prohibited & Restricted Plants & Fungi

where applicable, and that where such certification relies on third party audits, analysis, industry codes, or equivalence of international standards to demonstrate compliance, that certificates are current and available:

- 2) acknowledges that the Customer, and Supply Chain Customers of the Customer, will rely on the accuracy of the Product Information for food quality, safety and labelling purposes;
- 3) certifies that the accuracy of the Product Information is limited to the following degree: -
 - (a) that the Product Information in relation to ingredients obtained from a third party relies in good faith on Product Information provided by that third party;
 - (b) that the information is, to the best of the supplier's knowledge (having undertaken all reasonable verification procedures), true and accurate in relation to all other substances and processes;
- 4) agrees that all Product it supplies to the Customer will conform with the Product Information unless otherwise agreed to in writing and in advance by the Customer;
- 5) will immediately inform the Customer (and confirm in writing as soon as possible) if the supplier becomes aware of any error or omission in the Product Information;
- 6) will inform the Customer in writing and in advance of any change to the Product Information provided herein (including any changes that result from new or modified processes) if and when the supplier becomes aware of such changes; and
- 7) acknowledges that the Customer may provide the Product Information to
 - (a) regulatory agencies in relation to any matter raised by such agencies:
 - (b) courts and other legal tribunals for the purposes of any proceedings; and
 - (c) to its related businesses and partners who are involved in the acquisition, use, sale or compliance of the Product, under this same restriction as to disclosure.

but will otherwise NOT disclose the Product Information.

8) acknowledges that, subject to the prior written agreement of the supplier and any restrictions nominated by the supplier in regard to disclosure of confidential information, the Customer may provide the Product Information to its own customers subject to those customers ensuring the information is not further disclosed.

COMPANY NAME Signed for and on behalf of	Sun Pharmaceutical Industries (Australia) Pty Ltd
NAME (Please print)	Cassandra Amos
JOB TITLE (Please print)	Quality Assurance Officer
AUTHORISED SIGNATURE	cpmos.
DATE OF AUTHORISATION	08-April-2020

1.5 CUSTOMER DETAILS (WHERE KNOWN)

COMPANY NAME			
NUMBER / STREET / SUBURB			
CITY / COUNTRY / POST CODE			
CUSTOMER CONTACT NAME			
CUSTOMER'S PRODUCT NAME			
CUSTOMER'S PRODUCT CODE			
Cus	tomer Internal	Use Only	
Internal Product Code/Description			
Version No.			
Reason for Update			
Received and Reviewed By			
Approved [Yes/No]	_	Date:	
Signature:	Insert signatur	e here	

1.6 DEFINITIONS / REFERENCES

References to the "Code" or specific "Standards" throughout this document refer to the standards outlined in the Australia New Zealand Food Standards Code. The Australia New Zealand Food Standards Code can be viewed at: http://www.foodstandards.gov.au/foodstandardscode/

The AFGC provides some industry guides, specifically on how to apply date marking, and the AFGC Allergen Management and Labelling Guide which are available from the AFGC website: http://www.afgc.org.au/

Additional related documents on allergen management and VITAL (Voluntary Incidental Trace Allergen Labelling) documents can be viewed at: http://www.allergenbureau.net/vital/

1.7 CHECKLIST AND ATTACHMENTS

Page 2 has been signed and dated (Section 1.4)

Current Certificates attached - if applicable (Section 3.2.3 and Section 5.2)

Supplier C of C, or C of A for analysis - if applicable (Section 7)

Other associated documents attached as requested by the customer (e.g. MSDS, HACCP certification, product specification, and related documents)

1.8 Status of completion for each section:

COMPLETED Section 1 - Contact details and declaration **PARTIAL** Section 2 - Product Information & Ingredients COMPLETED Section 3 - Compositional information COMPLETED Section 4 - Foods requiring pre-market clearance **PARTIAL** Section 5 - Nutrients & consumer information claims **PARTIAL** Section 6 - Product shelf life, storage & packaging COMPLETED Section 7 - Chemical, microbial, organoleptic & physical specifications COMPLETED Section 8 - Additional comments

Check Box if help is needed identify mandatory sections of form which have NOT been completed:

2	PRODUCT INFOR	MATION & INGRED	DIENTS				
2.1 I	PRODUCT DESCRIPT	ION (Physical and techn	ological de	escription)			
Popp	y Seed (Kidney shaped	d seeds approimately 1.0	00 mm x 0	.75 mm)			
2.2 L	EGAL DESCRIPTION	/ SUGGESTED LABELI	LING DES	CRIPTION			
Popp	y Seed						
2.3	PRODUCT APPLICATION	ION AND INTENDED US	SE				
2.3.1	Specify the intended u	ise of the product					
	Food may be used a	s an ingredient, or may	be retail-	ready finished p	oroduct		
2.3.2	Specify which best de	scribes the product					
	Solid, semi-solid or p	powder substance, rea	dy for cor	nsumption			
							_
	COUNTRY OF ORIGIN				:- : : 4- 4	41-1	
2.4.1	Declaration:	opriate overarching coun	itry or origi Cour		ich applies to i	inis product :	
	Grown in	Aı	ıstralia	y.			
	0.0111111	Į At	iotrana				
2.4.2	Indicate if the lo	ocal content of ingredient	s/compon		om Australia exceeds 95%	yes Yes	s/No
2.4.3	Are the primary composite from more than one co	onents, from which this pountry?	product is I	made or derived,	sourced	No Yes	s/No
2.4.4	The IMPOR	g apply in determining co RTED COMPONENTS ha The PRODUCT h 0% or more of total product racteristic of the product	ave underg nas underg ict costs a	gone substantial f gone substantial f re incurred in the	transformation transformation country stated	No Yes	s/No s/No s/No s/No
2.5	COMPONENT TYPE						
Sp X	product is a single co	mponents present in pro component substance edients, which may inclu- arious ingredients which	de compo	ound substances			
2.6 II	NGREDIENT DECLAR	ATION					
Specify Compo	y all ingredients including foo ound substances must specif	d additives in descending order y all ingredients and additives the food additive name or coo	present and	the characterising in	gredient or compo	onent. Food addi	
	many components ar		`		OMPONENT		tch

onents are in this product?

Number of COMPONENTs do not match components listed in table

COMPONENT NAME	PERCENT OF TOTAL	COMPOUND SUBSTANCE INGREDIENTS Full breakdown list of components in compound ingredient including additive code numbers	Characterising component %
Poppy Seed	98-100%		
Extraneous Matter		not greater than 2.0% by weight	
Insects defiled		not greater than 1.00%	
Insect excreta		not greater than 2,500 mg//lb	
Excreta other		not greater than 3 mg/lb	

2.6 INGREDIENT DECLARATION INCLUDING PERCENTAGE LABELLING (continued)

2.6 INGREDIENT DECLARATION INCLUDING PERCENTAGE LABELLING (continued) COMPONENT NAME COMPONE			
COMPONENT NAME	PERCENT OF TOTAL	INCENT	
	%	including additive code numbers	component %
Poppy Seed	98-100%		
Extraneous Matter		not greater than 2.0% by weight	
Insects defiled		not greater than 1.00%	
Insect excreta		not greater than 2,500 mg//lb	
Excreta other		not greater than 3 mg/lb	
	<u> </u>		
	1		

2.7 PROCESSING AIDS

Specify all processing aids used in the manufacture of this product not otherwise declared in the ingredient list.

NAME OF PROCESSING AID	FSC ADDITIVE NUMBER OR EC (as applicable)	PERMITTED USE AND CLASS NAME

3 COMPOSITIONAL INFORMATION

3.1 MANDATORY ADVISORY OR WARNING STATEMENTS & DECLARATIONS

("Yes" response triggers a mandatory advisory or warning statement. Refer Standard 1.2.3 of the Code)

FOOD / COMPONENT	PRESENT YES / NO
Bee pollen presented as a food or ingredient	No
Propolis presented as a food or ingredient	No
Unpasteurised milk and unpasteurised liquid milk products	No
Aspartame or aspartame-acesulphame salt (or phenylalanine)	No
Unpasteurised egg products	No
Quinine	No
Kola beverages containing added caffeine	No
Guarana or extracts of guarana	No
Phytosterol esters	No
Tall oil phytosterols.	No
Cereal-based beverages, where these foods contain no more than 2.5% m/m fat and less than 3% m/m protein, or less than 3% m/m protein only.	No
Evaporated and dried products made from cereals, where these foods contain no more than 2.5% m/m fat and less than 3% m/m protein, or less than 3% m/m protein only, as reconstituted according to directions for direct consumption.	No
Milk, and beverages made from soy or cereals, where these foods contain no more than 2.5% m/m fat.	No
Evaporated milks, dried milks and equivalent products made from soy or cereals, where these foods contain no more than 2.5% m/m fat as reconstituted according to directions for direct consumption.	No
Royal jelly presented as a food or ingredient	No
Polyols, Isomalts, Polydextrose (Lactitol, Maltitol, Maltitol syrup, Mannitol, Xylitol, Erythritol, Isomalt, Polydextrose, Sorbitol)	No

3.2 ALLERGEN MANAGEMENT & CONT	ΓROL	Yes/No			
3.2.1 Does the facility have a Food Safety	Yes				
3.2.2 Does the facility have a documented	d allergen management plan?	Yes			
IF YES, does this include the manag	gement of cross contact allergens?	yes			
3.2.3 Has the Food Safety Program been	independently audited and certified	? Yes			
If Yes provide name of Certi	fying Body HACCP - Derek Wilson Au	dit Services			
Date of most recent audit /	inspection 24-July-2020	Provide copy of certificate			
3.2.4 Indicate if any of the following is app cross contact within the manufacturi		_			
validated cleaning procedures	produ	uction scheduling			
control of personnel movement i	n factory X staff	training			
X documented procedures and controls X isolated storage of allergens					
raw material sourcing & tracing X dedicated equipment					
other					

3.3 INGREDIENTS TO BE DECLARED AS ALLERGENS OR SULPHITE

Please insert **YES** or **NO** to indicate if the product contains, or was manufactured using, any ingredient, additive or processing aid which has been derived from the following food sources. Highly processed derivatives must always be declared. Carefully assess compound ingredients for hidden allergens. [** Lupin included as a possible future addition to the Food Standards Code.]

Yes/No

No	Cereals containing gluten & their products [wheat, rye, barley, oats, spelt]
No	Crustacea & crustacea products
No	Egg & egg products
No	Fish & fish products (including mollusc with or without shells and fish oils)
No	Lupin & Iupin products [** not a mandatory labelling allergen at this time]
No	Milk & milk products
No	Peanut & peanut products
No	Sesame seed & sesame seed products
No	Soybean & soybean products
No	Tree nuts & tree nut products
	Reserved for future allergen - left blank intentionally

No

Sulphites, present in ingredients, additives or processing aids

3.3.1 Complete all coloured rows corresponding with "YES" declaration provided above.

3.3.1 Complete all colou	SOURCE NAME The	DERIVATIVE NAME			PROCESS
ALLERGENIC	allergenic food from which	Ingredient, additive or	PROPOI	RTION (%)	Allergenic
SUBSTANCE	ingredient is derived (e.g. wheat)	processing aid (e.g. maltodextrin)	Derivative in product	Protein in derivative	protein is removed?
Cereals containing gluten					
and their products					
[wheat, rye, barley, oats,					
spelt & derived product					
e.g. wheat maltodextrin]					
Crustacea					
& crustacea products					
·					
Egg					
& egg products					
a ogg producto					
Fish					
& fish products					
(including mollusc extract					
and fish oils)					
Lunin					
Lupin & lupin products					
& lupin products					
Milk					
& milk products					
& milk products					
Peanut					
Realiut Repeanut products (including					
peanut oil)					
Sesame Seed					
& sesame seed products					
(including sesame oils)					
Soybean & soybean products					
(including soybean oils)					
(o.aa.iig ooyboaii olio)					
Tree nuts					1
& tree nut products					
a troo hat products					1
December					
Reserved for future					
allergen					

3.4 ALLERGEN CROSS CONTACT

Yes/No Yes

3.4.1 Except for any allergens listed in Section 3.3, does your company have on site and handle ANY OTHER allergenic substances listed below?

IF YES, complete ALL columns with respect to the potential cross contact allergens based on information received through YOUR supply chain AND YOUR manufacturing processes.

**Refer to VITAL procedure and decision tree.

http://www.allergenbureau.net/vital/

3.4.2 All columns must be completed WHERE HIGHLIGHTED

3.4.2 All columns i	nust be co	ompietea w	HERE HIGHLIGHTED)	
ALLERGENIC SUBSTANCE	PRESENT IN SAME FACILITY Yes/No	PRESENT ON SAME LINE Yes/No	SOURCE FOOD The allergenic food from which ingredient is derived (e.g. wheat)	DERIVATIVE NAME Ingredient, additive or processing aid (e.g. maltodextrin)	TOTAL PROTEIN** protein level by VITAL, or specify "particulate" mg/kg
Cereals containing gluten & their products	Yes	No	Barley		
Crustacea & crustacea products	No				
Egg & egg products	No				
Fish & fish products (inc mollusc & oils)	No				
Lupin & lupin products	No				
Milk & milk products	No				
Peanuts & peanut products (inc peanut oil)	No				
Sesame Seed & sesame products	No				
Soybeans & soybean products (inc soybean oil)	No				
Tree nuts & tree nut products	No				
Reserved for future allergen					

3.4.3 Is cross contact allergen present in particulate form in the facility or on same	e iines?
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Nο	Vec/No

3.4.5 Have cross contact allergen levels been assessed using the VITAL procedure?

IF NO, Provide appropriate precautionary statement for this product in box below:

No	Yes/No

Barley used in straw pellet mill so it does not block during shut down. Not stored near poppy seeds

3.5 INTERNATIONAL ALLERGEN, LABELLING & INFORMATION REQUIREMENTS

	/ COMPONENT	PRESENT	ING & INFORMATION REQUESTION NAME OF FOOD	DERIVATIVE NAME
FOOD,	COMPONENT	(Yes/No)	(e.g. apple)	(e.g. cider vinegar)
Gelatine	beef - collagen	No		
Gelatine	other source	No		
Seafood	Algae/carrageenan	No		
products	Shellfish (Mollusc)	No		
Fungi	Matsutake mushroom	No		
i ungi	Other mushroom	No		
	Avocado	No		
	Banana	No		
	Pome fruit - apples, pears	No		
Fruits	Stone fruit - cherry, peach, plum, apricot.	No		
	Berry Fruits - blueberry, kiwifruit, strawberry	No		
	Citrus Fruits - grapefruit, lemon, lime, orange	No		
Grains,	Buckwheat	No		
Seeds, Nuts &	Coconut, poppy, sunflower, etc	Yes	Poppy Seed	Papaver somniferum
Spices	Mustard	No		
	Tomato	No		
	Yam	No		
	Allium genus - chive, leek, onion, garlic, spring onion	No		
	Legumes -			
Vegetables	other than peanut soybeans & lupins	No		
	Umbelliferae - aniseed, carrot, celery, celeriac, chervil, cumin, dill, coriander, fennel, parsley, parsnip	No		
Yeast & Yeast Products (including yeast extracts) Tick box if hydrolysed or autolysed		No		
Herbs Tick box if herb / herb extract		No		
Spice (excluding mustard) Tick box if spice / spice extract		No		

3.6 ADDITIONAL LABELLING & INFORMATION REQUIREMENTS

3.6 ADDITIONAL LABELLING &		PRESENT		INFORMATIO	N
FOOD / COMPONENT		(Yes/No)	TO BE PROVIDED WHERE PROMPTED		MPTED
	Butylated hydroxyanisole (BHA)	No	amount added (milligram/kilogram)		
Antioxidants	Butylated hydroxytoluene (BHT)	No	amount added (milligram/kil	ogram)	
	Other antioxidants	No	Specify type:	.,	
Added Caffei			amount added (milligram/kil		
	ne ally occurring)	No	amount added (milligram/kil	ogram)	
Alcohol (Res	idual)	No		l % v/v:	
			specific gravity if product is a Specify types of	aiconoi:	
			fats and oils:	-lr 10	V /N -
	Animal	No	Has fatty acid composition been Specify the process used to alter		Yes/No
l <u>-</u> .			- CPOORLY THE PROCESS GOOD TO WHICH	composition:	
Added Fats & Oils			Specify types of		
			fats and oils: If Palm oil is present, is this RSP	O certified?	Yes/No
	Vegetable	No	Has fatty acid composition been		Yes/No
			Specify the process used to alter	composition:	
			Specify type of vegetable protein		
	Acid	No	Specify type of vegetable protein	•	
Hydrolysed Vegetable	Hydrolysed		100% hydrolysis		
Proteins	Enzyme Hydrolysed		Specify type of vegetable protein	:	
		No	100% hydrolysis	<u> </u>	
			Name of sweetener	Number	Amount (mg/kg)
Intense swee	etener	No			3, 3,
			Name of preservative	Number	Amount (mg/kg)
Preservatives	3	No			
			Name of flavour enhancer	Additive nu	umber
Flavour enha	ncers	no			
Added Colou	rs	No			
Added Flavours					
		No			
		140			
Added Salt		No	amount added (milligran		
Added Sugar		No	amount added (gran	n/100g)	

	List specific component:	Provide relevant details necessary for consumer advice:
THER		
O YN.		
۵ ۲		

3.7 QUARANTINE & IMPORT/EXPORT INFORMATION REQUIREMENTS			
FOOD / COMPONENT	PRESENT (Yes/No)	ADDITIONAL INFORMATION TO BE PROVIDED WHERE PROMPTED	
	(165/140)	Specify type of animals	ED WHERE PROMPTED
Animal & Animal products		Specify type of animal derivatives	
(e.g. animal flesh, organs, stock, gelatine, animal fat, tallow, milk,	No	Specify country/ies of origin	
collagen from skin and / or hides etc)		Describe any heat processing used in the manufacture of this product (temperature/time):	
		Specify type of animals (tick appropriate box)	
		Specify type of meat derivatives	
Meat & Meat products (e.g. animal flesh, animal organs,		Specify source of meat products (i.e. Country and city):	
meat extracts)	No	Describe any heat processing used in the manufacture of this product (temperature/time):	
		How do you ensure products are derived from animals free of bovine spongiform encephalopathy (BSE)?	
		Specify type of birds (tick appropriate box)	
Bird & Bird products		Specify type of bird derivatives	
(e.g. meat, fat, eggs, extracts, feathers, feet, etc.)	No	Specify source of bird products (i.e. Country and city):	
		Describe any heat processing used in the manufacture of this product (temperature/time):	
		Specify type of fish:	
Fish & Fish products		Specify type of fish derivatives	
(e.g. smoked salmon, pilchards, shark fin, fish roe, etc)	No	Specify source of fish products (i.e. Country and city):	
		Describe any heat processing used in the manufacture of this product (temperature/time):	
		Specify type of honey or honey derivatives	
Honey & Honey products	No	Specify source of honey products (i.e. Country and State):	
		Describe any heat processing used in the manufacture of this product (temperature/time):	

		CLEARANCE

- 4.1 NOVEL FOODS (F
 - (Refer Standard 1.5.1 of the Code)
- 4.1.1 Is this product (or any of its components) listed as a novel food in the standard?

No	Yes/N
NO	Yes/IN

4.2 QUARANTINE TREATMENTS

Specify if this product (or any of its components) has been treated with the following:

TREATMENT METHOD	USED ON ANY COMPONENT	SPECIFY TREATED INGREDIENT
Steam sterilisation	No	
Ionising (gamma) irradiation	No	
Ethylene oxide	No	
Other fumigants or sterilants	No	

4.3 FOOD PRODUCED USING GENE TECHNOLOGY (Standard 1.5.2)

	IF NO, specify which of the following are applicable:
	synthesis by GM micro-organisms, but with the exemption of use of GM feedstock?
	product that come from genetically modified (GM) plants or animals, or are the result of
4.3.1	Are there any ingredients (including food additives, processing aids and enzymes) in this

No	Yes/No
sence	

No GM varieties of this food / ingredient available
Non GM variety is used
Identity preservation program in place

Go	to	Quest	ion 4	.3.7	and (conti	nue

Analytical testing confirms absence					
Verifiable documentation of status					
Other – Specify					
-					

GM CROSS CONTAMINATION IN FOODS AND INGREDIENTS	Yes/No
4.3.7. Is this a raw/bulk commodity which is transported by freight/tanker AND where the freight/tanker could have previously been used to transport other GM product?	No
4.3.8. Is this product manufactured or stored at a production site where genetically	
modified protein or DNA is used for the manufacture of other products?	No
4.3.9. Is there an identity preservation system separating non GM and GM components	
to ensure the absence of genetically modified material in this product?	No
Specify details:	
4.3.10. Has Polymerase Chain Reaction (PCR) testing for GM materials been carried out?	No

4.3.11. Is any GM food or GM ingredient unintentionally present at MORE THAN 10g/kg

a information Form			F	rage	14					Print date
4.3.12. (OPTIONAL) Ar	e any ingre	diante d	ariyad from	n an	animal whi	ch has hoo	n fod wit	·h		No
feedstock contain									s?	NO
	iii ig Civi ii ig	rodiorito	or ingreat	Orrico	o donvou no		o organ		<u>. </u>	l
Specify details:										
5 NUTRIENT	S & CON	SUME	R INFO	₹M.	ATION CI	LAIMS				
5.1 NUTRITION INFOR										
- 4 4 50 15 15 15						4 5				
5.1.1 Please specify th					•	1.5 mm	gram			
5.1.2 For nutrition infor		•						graı		
Complete nutrient tab	le below. N	1andato	ry nutrients	hig	hlighted in l	blue and bo	lded, ot	hers	s optiona	al.
NUTRIENT		AVG	QUANTITY	1	% DI per	AVG QL	JANTITY			
HOTRIENT		PE	R SERVE		serve	per '	100 g			
Energy			#VALUE!	kJ	#VALUE!		1914	kJ	Martinia	
Protein, total			#VALUE!	g	#VALUE!		19.8	g		ent information
- Gluten		nc	t detected			not de	etected			SUPPLIE
Fat, total			#VALUE!	g	#VALUE!		32.5	g	AO	OUT LIL
- saturated			#VALUE!	g	#VALUE!		4.1	g		
- transfat									DO NO	OT leave bo
- polyunsaturated			#VALUE!	g			21.9	g		elds blank.
- monounsaturated			#VALUE!				6.5	g		ers, or text ' " with value
Cholesterol			#VALUE!				(II			/ailable" or
Carbohydrate			#VALUE!		#VALUE!		12	g		cted" for glu
- sugars			#VALUE!	g	#VALUE!		2.8	g		
Dietary fibre, total			#VALUE!	g	#VALUE!		21.6	g		
Sodium			#VALUE!	mg	#VALUE!		5.9	mg		
Potassium			#VALUE!	mg		7070 mg				
5.1.3 Additional nutrient	e - vitamine	miner	als and oth	er n	utritiva subs	stances			•	
Specify only one target							:			
, , , ,		•	XAd			ung Childr	_		Infants	j
VIITAMINIO			0/ DDI /		MINER					0/ DDL/
VITAMINS specify which vitamin	AVG QUA		% RDI / serve	er	MINERA becify which				NTITY	% RDI / serve
specify which vitairin	per 100	g	Serve	아	Decily Willeri	IIIIIIciais	per 10	00	g	Serve
NOTE: there is no perr	nission to F	ORTIF	foods with	h thi	s substance	e indicated	with **			
Insert any other nutrie	ent or biolo	gically	active sub	sta	nce					
NAME OF SUBSTANC					AVG QU	ANTITY per	100 g		%RDI/	serve
	Se	e Typic	al analysis							
5.1.4 Please provide the	ne following	analytic	al data:							
	% Ash		0%			Estimat			N/A	
9	6 Moisture	7-1	0%		acc	ounted for p	oer 100	g	N/A	
5.1.5 Please specify ho	w the carbo	ohydrate	e value has	be	en determin	ed:				
		,		'		_				

Difference as defined in Standard 1.2.8

Available Carbohydrate as defined in Standard 1.2.8

Other - specify: LTM158 Unknown

For laborator	Analytical – e.g. Laboratory y analysis, specify date of	Tested f analysi	is:	
	ABILITY TO MAKE CER' wif the product is suitable		in product intended for the following cons	Sumer uses
Орсси	SPECIFY IF SUITABLE		HOW HAS THIS BEEN VALIDATED?	CERTIFICATE AVAILABLE (Yes/No)
	Halal	yes		
	Kosher	Yes		Yes
	Organic	No		
	Biodynamic	No		
	Ovo-lacto-vegetarian	Yes	Suitable refer section 3.7	
	Lacto-vegetarian	Yes	Suitable refer section 3.7	
	Vegan	Yes	Suitable refer section 3.7	
A copy of re	levant certificates must	be pro	vided as attachments to form	
	PRODUCT SUITABILITY	FOR (es / No	SPECIFY PARTICULAR CLAIMS	HOW IS CLAIM VALIDATED?
	"Free" claims	No		
	Sustainability claims	No		
	Llumana traatmast	No		
	Humane treatment	No		
	Any other claims			
	Any other dains			

DURABILITY, PACKAGING AND SUPPLY CHAIN

6.1 SHELF LIFE

6.1.1 Please complete the following details:

	PRODUCT As unopened pack of		PRODUCT - ONCE IN USE resealable pack or bulk container		
Specify shelf life	2	Years	·		
Temperature control during storage	Is required ?	No	Is required ?		
			Specify range:		
Temperature control	Is required?	No			
during transport					
Specify any OTHER storage requirements:	Cool Dry Storage				

6.1.2 Specify the type of date mark to be used: **Best before**Refer to AFGC Date Marking Guide

6.2 POTENTIAL HAZARDS

6.2.1 Are there any potential hazards associated with the product?

No	Yes/No

6.3 TRANSPORT

How is product transported and packaged?

Packaged for catering/manufacturing supply

^ 4 TD			\sim 1 to	
64 TR	$\Delta I I \vdash$	$M \vdash \Delta$	< I I I I	EMENT

1.1	1	Specify	/ which	method o	f trade	measurement is used:	
ł. 1	1	Specify	/ wnich	metnoa o	it trade	measureme	nt is usea:

6.4.2 What is the package size 15.00

6.4.3 Target Fill (if applicable)

6.4.4 Drained Weight (if applicable)

6.4.5 IF AQS is used, what is the statistical variance in the fill measurement?

	net quantity	
kg	(specify unit of	measure)
	(specify unit of	measure)
	(specify unit of	measure)
		ı

6.5 TRACEABILITY

Please provide any general comments about the traceability coding used on the product:

Each pallet has own unique lot code. Pallet configuration 9 high, 6 per row. 54 bags per pallet

Please specify the following where applicable:

TRACKING CODE	UNIT				SHIPPER (if applicable)			
Type of Primary Coding		Date code		Batch number		Date code		Batch number
(Please TICK as appropriate)		Product code	X	Lot number		Product code		Lot number
Method of coding	Printing on side of bag							
Location of code	Left hand side gusset							
Number of characters in code	8.00							
Example of coding format	ormat L2001507							
Coding translation	20=year remaining 5 numbers = pallet number							

6.6 PRODUCT PACKAGING

- 6.6.1 Are tamper evident controls included in the packaging design?
- 6.6.2 Has unit packaging been assessed for migration of substances into food?
- 6.6.3 Are engineered nanoparticles present in unit packaging?

Yes	Yes/No
No	Yes/No
No	Yes/No

6.6.4 Are you a signatory to relevant packaging stewardship in Australian or NZ?

No Yes/No

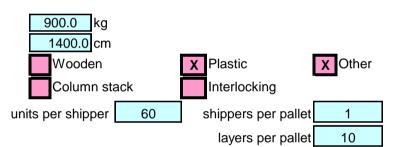
6.6.5 Provide a general description of unit packaging:

666	Complete the following table for questions related to packaging of unit package and/or shipper

	PACKAGING	UNIT	SHIPPER
Туре	Packaging format	3 ply	
	Ceramic	No	
	Glass	No	
Specify	Metal	No	
components /	Paper / cardboard	Yes	
material used	Packing materials	Yes	
in packaging	Plastics	No	
	% of total using recycled component	0%	
Seal	What is the seal method?	Sewn	
	Height (mm)	660	700
Dimensions	Width (mm)	440	480
	Depth (mm)	110	130

6.7 PALLET CONFIGURATION

- 6.7.1 Gross weight of loaded pallet
- 6.7.2 Stack height of loaded pallet
- 6.7.3 Specify the type of pallet
- 6.7.4 What is the pallet pattern
- 6.7.5 Number of:



7 SPECIFICATIONS FOR COMPLIANCE

Test Methods are mandatory and must quote AOAC methods or recognised independent Australian or International standards. Where a supplier's internal test method is quoted a copy must be attached. Also state if Certificate of Analysis (C of A) or Certificate of Conformance (C of C) can be provided.

7.1 ORGANOLEPTIC SPECIFICATIONS

(Examples may include flavour, colour, aroma, texture etc)

			AVAILABILITY		
TEST / PARAMETER	SPECIFICATION	TEST METHOD	C of A	C of C	
colour	Blue / Grey	Visual - AML001			

7.2 PHYSICAL SPECIFICATIONS

(Examples may include particle size, shape, specific gravity, metal detection, foreign matter tolerances, physical defect tolerances etc as appropriate for the product)

, , , , , , , , , , , , , , , , , , , ,			AVAILA	BILITY
TEST / PARAMETER	SPECIFICATION	TEST METHOD	C of A	C of C
Purity	98 - 100 %	AML001	Yes	
,				

7.3 MICROBIOLOGICAL SPECIFICATIONS

(Examples may include standard plate count, yeasts & moulds, coliforms, salmonella, listeria etc)

			AVAILA	BILITY
TEST / PARAMETER	SPECIFICATION	TEST METHOD	C of A	C of C
E Coli	< 3 CFU/g	M52		

7.4 CHEMICAL SPECIFICATIONS

(Examples may include pesticide residue screen, antibiotic residue screen, heavy metal screen, aflatoxins screen, salt, acid, pH, moisture, brix, Aw, Nutrition Information Test, etc)

	i, moistare, brix, Aw, Nathtiori informa	, ,	AVAILA	BILITY
TEST / PARAMETER	SPECIFICATION	TEST METHOD	C of A	C of C
See Typical analysis				Yes

	8.1 Do you have any comments or additional information? No Yes/No				
Question Number	Line Number	Comments			

8.2 ADDITIONAL MANUFACTURING SITE INFORMATION (if required)

When nominating product is supplied from more than one manufacturing site, the details provided must be applicable to product coming from any of the sites. For example, if particular allergens occur at only one site then the information provided in the form should identify that the allergens are present even though batches of product made at other sites may be allergen free.

	0 1	, ,		
	COMPANY NAME			
SITE: #	#4 NUMBER / STREET / SUBURB			
	STATE / COUNTRY / POST CODE			
	COMPANY NAME			
SITE: #	#5 NUMBER / STREET / SUBURB			
	STATE / COUNTRY / POST CODE			
	COMPANY NAME			
SITE: #	#6 NUMBER / STREET / SUBURB			

STATE / COUNTRY / POST CODE