9 - 11 Antill Street, Yennora, New South Wales 2161, Australia
CERTIFICATE OF ANALYSIS

Product:

Super Feast Jing HS Capsules

Sponsor:

Superfeast (JEMS Superfoods)

Batch No .:

1765a

Pack Size:

Bulk

Units Produced:

36000 capsules

Date of Manufacture:

Oct 2023

Expiry Date:

Sponsor's responsibility when stored at below 30°C and protected from light.

Dosage Form:

Capsule, Hard Shell

TEST	SPECIFICATION	RESULT
Appearance	A clear two- piece size "00" veggie capsule containing a brown -coloured powder	Conforms
Average Fill Weight	514.5mg ± 7.5% w/w (475.9 to 553.1 mg)	542.7mg
Average Total Weight	634.5 mg ± 7.5% w/w (586.9 to 682.1mg)	662.7mg
Cordyceps Extract Powder 10:1	0.111g/g	
Equivalent to mycelium	1.11g/g (90% to 120% of stated amount), not assayed, Quantified by Input	QBI
Lycium barbarum (Goji berry) Extract Powder 10:1	0.111g/g	QBI
Equivalent to fruit	1.11/g (90% to 120% of stated amount), not assayed, Quantified by Input	
Eucommia ulmoides Extract Powder	0.111g/g	QBI
Equivalent to stem barks	1.11g/g (90% to 120% of stated amount), not assayed, Quantified by Input	
Rehmannia glutinosa Extract Powder 10:1	0.0555g/g	QBI
Equivalent to dry root	0.555/g(90% to 120% of stated amount), not assayed, Quantified by Input	
Cistanche salsa Extract Powder 10:1	0.0555g/g	QBI
Equivalent to stem	0.555g/g(90% to 120% of stated amount), not assayed, Quantified by Input	
Asparagus cochinchinensis Extract Powder 10:1	0.0555g/g	QBI
Equivalent to dry root	0.555g/g (90% to 120% of stated amount), not assayed, Quantified by Input	
Pesticide Residue	Meet Eur.Ph (2.8.13)	Rotational
Aflatoxins (subtotal of B1, B2, G1, G2)	20.0ppb maximum	Rotational
Heavy Metals	10.0ppm maximum	Rotational



Disintegration Time	30 minutes maximum @ 37.0°C (water)	Complies
Average Capsule Length 23.30mm±0.76mm		23.32mm
Total Aerobic Microbial	10 ⁴ cfu/g maximum	Complies
Total Yeast and Mould	10 ² cfu/g maximum	Complies
Bile-tolerant Gram- Negative Bacteria	10 ² cfu/g maximum	Complies
Escherichia coli	Not detected/g	Complies
Salmonella	Not detected/10g	Complies
Staphylococcus aureus	Not detected/g	Complies

Residual solvents: All raw materials in this product have been assessed and verified to be compliant with Ph Eur 5.4, and no solvents are used during the manufacturing process, so the further testing is not required.

The first 3 batches of finished products will be tested and then as part of a rotational testing program 1 batch in every 10 batches manufactured or annually, whichever comes first, will be tested thereafter.

QBI: Quantified by Input

Prepared by:	A	Date: 13 11/10 17
	QA Supervisor	
Reviewed by:	2	Date: 13/12/D
	Operations Manager	



CERTIFICATE OF CONDITIONAL RELEASE FOR FURTHER

PROCESSING

Product:

Superfeast Jing HS Capsules

Product Code:

BG10064

Batch No .:

1765a

Units produced:

36000

Date of Manufacture: Oct 2023

This document is to certify that the bulk product has been produced and controlled according to Good Manufacturing Practices and the requirements of the Marketing Authorisation and is, therefore, Released for further processing.

The documentation review process has involved the following elements.

- 1. The assurance that the operations under our control have been carried out in accordance with the requirements of the Australian code of Good Manufacturing Practice for Therapeutic Goods.
- 2. That a document review has been performed as required by the Document Checklist as well as the Release Checklist, which is incorporated into the batch documentation.
- 3. That the reconciliations of the yields of the bulk product are within approved limits and are acceptable.
- 4. That any reconciliations of the packaging components are within approved limits and are acceptable.
- 5. That the document package is complete and free from any known errors. That any omissions, anomalies, and deviations have been documented, explained, and appropriately authorised.
- 6. The recommended storage condition is consistent with the shelf life of the packed product. The bulk product is recommended to store no more than 3 months from date of manufacturing as the storage period for bulk without shelf-life justification before final packing. Sponsor is responsible for justification of the shelf life above 3 months for bulk as well as the shelf life of finished packed product.
- 7. That a formalised Labceuticals Pty Ltd Agreement is in place.
- 8. That the product described above complies with the relevant Marketing Authorisation.

Labceuticals Pty Ltd is a current holder of Therapeutic Goods Administration Manufacturing License: MI-2016-LI-11619-1

QA Supervisor

Signed



9 - 11 Antill Street, Yennora, New South Wales 2161, Australia TGA GMP Licence No: MI-2016-LI-11619-1

CERTIFICATE OF RELEASE FOR SUPPLY

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Super Feast Jing HS Capsules 240's

Product Code:

FP20097

Batch No.:

1766

Units produced:

247

Date of Packaging:

Nov 2023

This document is to certify that the product has been produced and controlled according to Good Manufacturing Practices and the requirements of the Marketing Authorisation and is, therefore, **Released for Supply.**

The documentation review process has involved the following elements.

- 1. The assurance that the operations under our control have been carried out in accordance with the requirements of the Australian code of Good Manufacturing Practice for Therapeutic Goods.
- 2. That a document review has been performed as required by the Document Checklist as well as the Release Checklist, which is incorporated into the batch documentation.
- 3. That the reconciliations of the yields of the finished product are within approved limits and are acceptable.
- 4. That any reconciliations of the packaging components are within approved limits and are acceptable.
- 5. That any reconciliations of the labels and or pre-printed packaging components are within approved limits and are acceptable.
- 6. That the document package is complete and free from any known errors. That any omissions, anomalies, and deviations have been documented, explained, and appropriately authorised.
- 7. That a formalised Labceuticals Pty Ltd Agreement is in place.
- 8. That the product described above complies with the relevant Marketing Authorisation.

Jessie Liu	A	13/12/2027
QA Supervisor	Signature	Date

Product:

Super Feast Jing HS Capsules 240's

Sponsor:

Superfeast (JEMS Superfoods)

Batch No .:

1766

Pack Size:

240 / Bag

Units Produced:

247units

Date of Manufacture:

Oct 2023

Expiry Date:

Oct 2025 when stored at below 30°C and protected from light.

Dosage Form:

Capsule, Hard Shell

TEST	SPECIFICATION	RESULT	
Appearance	A clear two- piece size "00" veggie capsule containing a brown -coloured powder	Conforms	
Average Fill Weight	514.5mg ± 7.5% w/w (475.9 to 553.1 mg)	542.7mg	
Average Total Weight	634.5 mg ± 7.5% w/w (586.9 to 682.1mg)	662.7mg	
Cordyceps Extract Powder 10:1	0.111g/g		
Equivalent to mycelium	1.11g/g (90% to 120% of stated amount), not assayed, Quantified by Input	QBI	
Lycium barbarum (Goji berry) Extract Powder 10:1	0.111g/g	QBI	
Equivalent to fruit	1.11/g (90% to 120% of stated amount), not assayed, Quantified by Input		
Eucommia ulmoides Extract Powder 10:1	0.111g/g	QBI	
Equivalent to stem barks	1.11g/g (90% to 120% of stated amount), not assayed, Quantified by Input		
Rehmannia glutinosa Extract Powder 10:1	0.0555g/g	QBI	
Equivalent to dry root	0.555/g(90% to 120% of stated amount), not assayed, Quantified by Input		
Cistanche salsa Extract Powder 10:1	0.0555g/g	QBI	
Equivalent to stem	0.555g/g(90% to 120% of stated amount), not assayed, Quantified by Input		
Asparagus cochinchinensis Extract Powder 10:1	0.0555g/g	QBI	
Equivalent to dry root	0.555g/g (90% to 120% of stated amount), not assayed, Quantified by Input		
Pesticide Residue	Meet Eur.Ph (2.8.13)	Rotational	
Aflatoxins (subtotal of B1, B2, G1, G2)	20.0ppb maximum	Rotational	
Heavy Metals	10.0ppm maximum	Rotational	



Disintegration Time	30 minutes maximum @ 37.0°C (water)	Complies
Average Capsule Length	23.30mm±0.76mm	23.32mm
Total Aerobic Microbial	10 ⁴ cfu/g maximum	Complies
Total Yeast and Mould	10 ² cfu/g maximum	Complies
Bile-tolerant Gram- Negative Bacteria	10 ² cfu/g maximum	Complies
Escherichia coli	Not detected/g	Complies
Salmonella	Not detected/10g	Complies
Staphylococcus aureus	Not detected/g	Complies

Residual solvents: All raw materials in this product have been assessed and verified to be compliant with Ph Eur 5.4, and no solvents are used during the manufacturing process, so the further testing is not required.

The first 3 batches of finished products will be tested and then as part of a rotational testing program 1 batch in every 10 batches manufactured or annually, whichever comes first, will be tested thereafter.

QBI: Quantified by Input

Prepared by:	QA Supervisor	Date:	13/11/23
Reviewed by:	Operations Manager	Date:	13/12/23



9 - 11 Antill Street, Yennora, New South Wales 2161, Australia CERTIFICATE OF ANALYSIS

Product:

Super Feast Jing HS Capsules 120's

Sponsor:

Superfeast (JEMS Superfoods)

Batch No .:

1767

Pack Size:

120 / Bottle

Units Produced:

2547units

Date of Manufacture:

Nov 2023

Expiry Date:

Oct 2025 when stored at below 30°C and protected from light.

Dosage Form:

Capsule, Hard Shell

TEST	SPECIFICATION	RESULT
Appearance	A clear two- piece size "00" veggie capsule containing a brown -coloured powder	Conforms
Average Fill Weight	514.5mg ± 7.5% w/w (475.9 to 553.1 mg)	539.9mg
Average Total Weight	634.5 mg ± 7.5% w/w (586.9 to 682.1mg)	659.9mg
Cordyceps Extract Powder 10:1	0.111g/g	303.31118
Equivalent to mycelium	1.11g/g (90% to 120% of stated amount), not assayed, Quantified by Input	QBI
Lycium barbarum (Goji berry) Extract Powder 10:1	0.111g/g	QBI
Equivalent to fruit	1.11/g (90% to 120% of stated amount), not assayed, Quantified by Input	
Eucommia ulmoides Extract Powder 10:1	0.111g/g	QBI
Equivalent to stem barks	1.11g/g (90% to 120% of stated amount), not assayed, Quantified by Input	
Rehmannia glutinosa Extract Powder 10:1	0.0555g/g	QBI
Equivalent to dry root	0.555/g(90% to 120% of stated amount), not assayed, Quantified by Input	
Cistanche salsa Extract Powder 10:1	0.0555g/g	QBI
Equivalent to stem	0.555g/g(90% to 120% of stated amount), not assayed, Quantified by Input	
Asparagus cochinchinensis Extract Powder 10:1	0.0555g/g	QBI
Equivalent to dry root	0.555g/g (90% to 120% of stated amount), not assayed, Quantified by Input	
esticide Residue	Meet Eur.Ph (2.8.13)	Complies
flatoxins (subtotal of B1, B2, G1, G2)	20.0ppb maximum	Complies
leavy Metals	10.0ppm maximum	Complies



rn'	30 minutes maximum @ 37.0°C (water)	Complies
Disintegration Time		23.17mm
Average Capsule Length	23.30mm±0.76mm	23.1711111
Total Aerobic Microbial	10 ⁴ cfu/g maximum	Complies
Total Yeast and Mould	10 ² cfu/g maximum	Complies
Bile-tolerant Gram-	10 ² cfu/g maximum	Complies
Negative Bacteria		Complies
Escherichia coli	Not detected/g	Complies
Salmonella	Not detected/10g	*
	Not detected/g	Complies
Staphylococcus aureus	to into in this product have been assessed a	nd verified to be compliant

Residual solvents: All raw materials in this product have been assessed and verified to be compliant with Ph Eur 5.4, and no solvents are used during the manufacturing process, so the further testing is not required

The first 3 batches of finished products will be tested and then as part of a rotational testing program 1 batch in every 10 batches manufactured or annually, whichever comes first, will be tested thereafter.

QBI: Quantified by Input

Prepared by:	QA Supervisor	Date:	13/12/20
	QA Supervisor		

Operations Manager



9 - 11 Antill Street, Yennora, New South Wales 2161, Australia TGA GMP Licence No: MI-2016-LI-11619-1

CERTIFICATE OF RELEASE FOR SUPPLY

Product:

Super Feast Jing HS Capsules 120's

Product Code:

FP20090

Batch No.:

1767

Units produced:

2547

Date of Packaging:

Nov 2023

This document is to certify that the product has been produced and controlled according to Good Manufacturing Practices and the requirements of the Marketing Authorisation and is, therefore, **Released for Supply.**

The documentation review process has involved the following elements.

- 1. The assurance that the operations under our control have been carried out in accordance with the requirements of the Australian code of Good Manufacturing Practice for Therapeutic Goods.
- 2. That a document review has been performed as required by the Document Checklist as well as the Release Checklist, which is incorporated into the batch documentation.
- 3. That the reconciliations of the yields of the finished product are within approved limits and are acceptable.
- 4. That any reconciliations of the packaging components are within approved limits and are acceptable.
- 5. That any reconciliations of the labels and or pre-printed packaging components are within approved limits and are acceptable.
- 6. That the document package is complete and free from any known errors. That any omissions, anomalies, and deviations have been documented, explained, and appropriately authorised.
- 7. That a formalised Labceuticals Pty Ltd Agreement is in place.
- 8. That the product described above complies with the relevant Marketing Authorisation.

Jessie Liu QA Supervisor

Signature

Date