

Japan Food Research Laboratories

Accredited by the Japanese Government

52-1 Motoyoyogi-cho, Shibuya-ku, Tokyo 151-0062, Japan http://www.jfrl.or.jp/

No. 16017889001-0301

1/1

Date issued: February 29, 2016

CERTIFICATE OF ANALYSIS

Client:

Showa Denko K. K.

5-1 Ohgi-machi, Kawasaki-ku, Kawasaki-shi, Kanagawa 210-0867, Japan

Sample name:

GLYCINE 160211A

Received date:

February 17, 2016

This is to certify that the following result(s) have been obtained from our analysis on the above-mentioned sample(s) submitted by the client.

Test Result(s)

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Test Item	Result	QL	N	M
Glycine			1	
Description	Conformable	(**************************************		
Identification (1)	Conformable			
Identification (2)	Conformable			
Purity		******		
Clarity and color of solution	Conformable	(20200)		
рН	Conformable (pH6. 2)			
Chloride	Conformable	Section 19		
Heavy metals	Conformable	(10000000)		
Arsenic	Conformable			
Loss on drying	Conformable (0.06 %)			
Residue on ignition	Conformable(Not more than	(man)		
	0.05 %)			
Assay (Content)	Conformable (99.9 %)			

QL: Quantitation limit N: Notes M: Method

Notes

1:"Food Additives," Notification No. 370 (1959) "Specifications and Standards for Foods, Food Additives, etc.," issued by the Ministry of Health and Welfare.

Signed for and on behalf of JFRL

Takeko Arai

Section of Analysis Documentation

Feb. 29, 2016
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Test Result(s)

Test Item	Result	QL	N M
Aerobic plate count	Less than 300/g	(*************************************	1
Coliform bacteria	Negative/1g		1
Escherichia coli	Negative/1g		1
Total combined molds and yeasts count	Less than 30/g		1
Coagulase positive staphylococci	Negative/0.01g	:	1
Salmonella	Negative/25g	******	2
Pseudomonas aeruginosa	Negative/1.1g		2
Clostridia	Negative/1g		3 3
Mesophilic aerobic spores	Less than 30/g	•••••	1
Gas-producing anaerobes	Negative	*******	1

QL: Quantitation limit N: Notes M: Method

Notes

- 1:The test method was specified by the client.
- 2:The Methods for the Microbiological Examination of Foods, NIHSJ-01-ST4:2009 (091014) "Detection method for Salmonella."
- 3:Heat-shocked conditions: 70 °C for 20 minutes.

Method

1:Surface spread plating method

2:Enrichment culture method

3:Clostridia Count Agar anaerobic plating method

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