



SIRIM QAS International Sdn.Bhd. (Company No.: 410334-X)
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TEST REPORT

REPORT NO. : 2017CE0175

PAGE : 1 OF 6

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Applicant : LAMACO SYSTEM SDN BHD
407 & 408, Jalan Perusahaan 6,
Taman Bandar Baru Mergong,
05150 Alor Setar, Kedah.

Product : INJECT SEAL ADVANCE

Reference standard /
Method of test : BS 6920-1: 2014
Suitability of non-metallic materials and products for use in contact
with water intended for human consumption with regard to their effect
on the quality of the water
Part 1: Specification

Description of sample : Received one (1) sample of **INJECT SEAL ADVANCE** for testing.
The sample was identified as:
BRAND : LaMaCO


Date received of
Complete Application : 14th December 2016

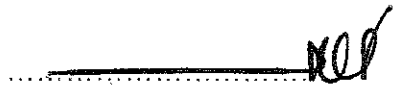
Job No. : J20161401626

Description of test
results : The test results of the submitted sample are described in Page 2 to
Page 6 of this test report.

Issued date : 23rd February 2017

Approved Signatory :


(NOR AZLIN BT M NOOR ARZMI, MMIC)
IKM M/3012/5811/2010
Testing Executive


(HAHNAS MAHBUT)
Head
Chemical & Consumer Section
Testing Services Department

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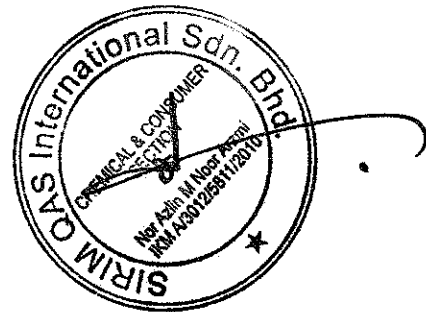
TEST RESULTS

TEST : ODOUR AND FLAVOUR OF WATER
PRODUCT : INJECT SEAL ADVANCE
BRAND : LaMaCO

No.	Type of Tests	Requirement under BS 6920-1: 2014 : Clause 4	Test results after final extraction	
			Unchlorinated extracts	Chlorinated extracts
1.	Odour	No discernable odour	Plastic	Plastic
2.	Flavour	No discernable flavour	N/A	N/A

- Note:**
1. The final unchlorinated and chlorinated extracts for odour tests were after 7th extractions
 2. Flavour test for unchlorinated and chlorinated extracts was not proceed since odour was detected
 3. N/A = not applicable

Comment: The sample tested did not comply with the requirements of BS 6920-1: 2014: Clause 4



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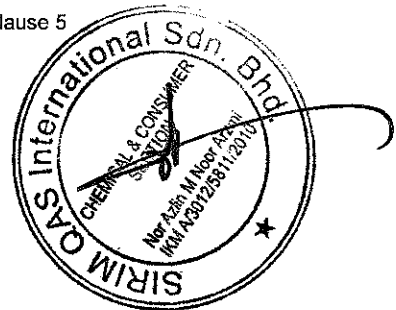
TEST RESULTS

TEST : APPEARANCE OF WATER
 PRODUCT : INJECT SEAL ADVANCE
 BRAND : LaMaCO

No.	Type of Tests	Methods Used	Requirement under BS 6920-1: 2014 : Clause 5	Test results after final extraction		
				Control	Sample	Net Increase
1.	Colour, mg/L Pt	APHA 2120 B	The increase in colour and turbidity of the water in the final extract shall not be more than 5 mg/L Pt or 0.5 FNU respectively	Less than 5	Less than 5	Less than 5
2.	Turbidity, FNU	APHA 2130 B		0.05	0.20	0.15

Note : The final extract for colour and turbidity tests were after 24 hours (1st extraction)

Comment : The sample tested complied with the requirements of BS 6920-1: 2014: Clause 5



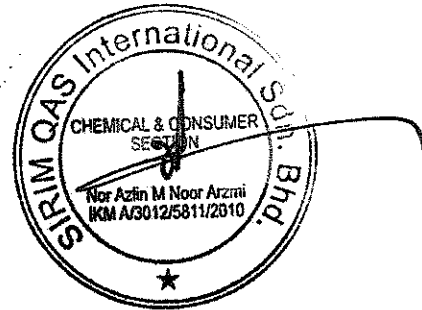
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TEST RESULTS

TEST : GROWTH OF AQUATIC MICROORGANISMS
 PRODUCT : INJECT SEAL ADVANCE
 BRAND : LaMaCO

Type of Tests	Requirement under BS 6920-1: 2014 : Clause 6	Test results after final extraction		
		After 5 th week	After 6 th week	After 7 th week
The mean dissolved oxygen difference (MDOD), mg/L	2.39 or less	0.11	0.08	0.07

Comment : The sample tested complied with the requirements of BS 6920-1: 2014: Clause 6

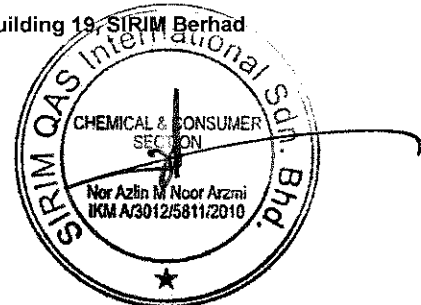


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TEST RESULTS

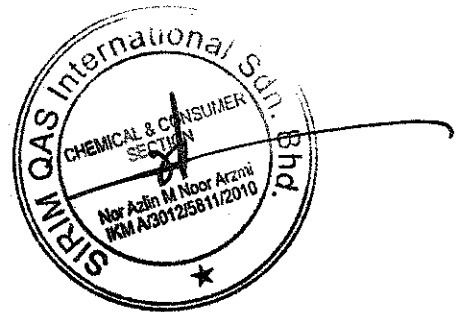
TEST ⁽¹⁾ : **EXTRACTION OF SUBSTANCES THAT MAY BE OF CONCERN TO PUBLIC HEALTH**
PRODUCT : **INJECT SEAL ADVANCE**
BRAND : **LaMaCO**
TEST RESULT : **REFER APPENDIX 1 – REPORT NO R001/17/B19/01**

Note : ⁽¹⁾ Test subcontracted to Industrial Biotechnology Research Centre, Building 19, SIRIM Berhad



Report No. : 2017CE0175

APPENDIX 1





TEST REPORT

REPORT NO: R001/17/B19/01

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Applicant : SIRIM QAS International Sdn. Bhd.
Chemical and Consumer Section, Building 16,
SIRIM Berhad.
(Pn Nor Azlin M. Nor Arzmi)

Manufacturer / Company : Lamaco System Sdn. Bhd

Sample/Trade Name : Inject Seal Advance

Reference Standard / Method of Test : 1. BS 6920-2.1:2014. Suitability of non-metallic materials and products for use in contact with water intended for human consumption with regard to their effect on the quality of the water
- Part 2: Methods of test - Section 2.1: Samples for testing.
2. BS 6920-2.5:2000+A2:2014. Suitability of non-metallic products for use in contact with water intended for human consumption with regard to their effect on the quality of the water
- Part 2: Methods of test - Section 2.5: The extraction of substances that may be of concern to public health

Description of Sample : Received one sample in good condition consisting of 3 pieces for testing with the following identification:

1. Model/Brand/Marking: Lamaco
2. Product Marking: Not provided
3. Reference No: J20161401626
4. Designation: Not provided
5. Date of Manufacture: Not provided
6. Nature of Material: Not provided
7. Storage Condition: Room temperature
8. Colour: Light Yellow
9. Shape/Form: Square
10. Dimension: Approximate length 54mm, width 54mm, thickness 10mm
11. Appearance: Solid Rough Surface
12. Opacity: Opaque

Date Received : 29 December 2016

Job No. : J001/17

Issue Date : 20 JAN 2017



TEST REPORT

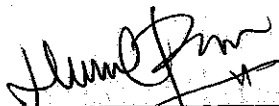
REPORT NO: R002/17/B19/01

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KEY PERSONNEL PARTICIPATING IN THIS TESTING

We, the undersigned, declare that the methods, results and data contained in this report faithfully reflect the procedures used and raw data collected throughout the testing.



(DR. NURUL IZZA NORDIN)
Deputy Technical Manager
Industrial Biotechnology Research Centre
SIRIM Berhad

20 JAN 2017

Date



(JUANI MAZMIN HUSIN)
Officer In-Charge
Industrial Biotechnology Research Centre
SIRIM Berhad

20 JAN 2017

Date



TEST REPORT

REPORT NO: R002/17/B19/01

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1.0 Test timetable

Receipt of sample: 29 December 2016
Maintenance of cell culture: 12 January – 13 January 2017
Extraction procedure: 12 January – 13 January 2017
Growth procedure: 13 January – 14 January 2017
End of test: 14 January 2017

2.0 Test method

2.1 Test summary

A screening procedure (simple cytotoxicity test) to test leachates from the **Inject Seal Advance** for biologically active compound was carried out according to BS 6920. Leachates from the sample after a 24-hour extraction at (23 ± 2) °C was used to prepare growth medium. The morphology of a mammalian cell line following a 24-hour culture in the growth medium was observed. **NO OTHER TESTS WERE UNDERTAKEN ON THIS PRODUCT.**

2.2 Significance and rationale

This method is only an initial screening test for substances potentially hazardous to health and suitable for all non-metallic materials that may be used in contact with water intended for human consumption. A satisfactory result indicates that the leachate probably does not contain significant amounts of acutely toxic substances, but it does not indicate the absence of small quantities of substances, which may be harmful on prolonged exposure.

2.3 Cell culture

American Type Culture Collection CCL-1, NCTC clone 929 Areolar Fibroblast Mouse.

2.4 Test procedure

The procedure was divided into three stages as follows.

2.4.1 Cell culture maintenance


Cells were grown in tissue culture grade flasks and routinely examined to ensure they remain healthy.

2.4.2 Sample extraction and preparation of growth media

Two sample pieces was selected. The sample surface area was measured at 5500 mm² to 9500 mm². The sample was extracted in 500 mL of reverse osmosis water at (23 ± 2) °C for 24 hours. A validation solution (800 mg/L zinc sulfate) and a blank were included in the test. Portions of the sample extracts, validation solution and blank were used to dilute the concentrated growth medium. Freshly trypsinized cells was added to each preparation and transferred into a 24-well tissue culture plate. The extracts were assessed in replicates of three. The plate was incubated at (37 ± 1) °C in a humidified atmosphere of 5 % carbon dioxide and 95 % air for 24 hours.

2.4.3 Effect on cell culture

The condition of cells in each well was examined microscopically. The presence or absence of a confluent cell layer, the presence of any irregular shaped cell or cells showing signs or 'rounding off', and the appearance of any cells floating in the growth medium was recorded.


JUANI MAZMIN BINTI HUSIN
Researcher
Bioprocess Programme
Industrial Biotechnology Research Centre
SIRIM Berhad



MS ISO/IEC 17025
TESTING
SIRIM NO.676

TEST REPORT

REPORT NO: R002/17/B19/01

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3.0 Results

The table below shows cytotoxicity effects of **Inject Seal Advance**


	Replicates	Effects on cell culture
Extract of sample piece 1	A	Healthy confluent monolayer
	B	Healthy confluent monolayer
	C	Healthy confluent monolayer
Extract of Sample piece 2	A	Healthy confluent monolayer
	B	Healthy confluent monolayer
	C	Healthy confluent monolayer
Validation solution	A	Cells 'rounding off' and floating. No monolayer
	B	Cells 'rounding off' and floating. No monolayer
	C	Cells 'rounding off' and floating. No monolayer
Blank	A	Healthy confluent monolayer
	B	Healthy confluent monolayer
	C	Healthy confluent monolayer

4.0 Analysis and interpretation

Cell cultures in the extract of samples showed healthy confluent monolayer, comparable to the blank, which indicate a non-cytotoxic response. Monolayer of cell culture was not present in the validation solution. The cells showed severe 'rounding off' and floating in the validation solution preparation, which indicate a cytotoxic response.

5.0 Conclusion

The sample **Inject Seal Advance** exhibited no cytotoxicity response under the conditions of this test.


JUANI MAZMIN BINTI HUSIN
Researcher
Bioprocess Programme
Industrial Biotechnology Research Centre
SIRIM Berhad



MS ISO/IEC 17025
TESTING
SAMM NO. 078

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TEST RESULTS

TEST : EXTRACTION OF METALS
PRODUCT : INJECT SEAL ADVANCE
BRAND : LaMaCO

No.	Type of Tests	Method Used	Requirement under BS 6920-1: 2014: Clause 8 (maximum allowed)	Test results after 24 hours extraction/final extraction ⁽¹⁾
1.	Aluminium (as Al), µg/L	APHA 3120 B	200	5
2.	Antimony (as Sb), µg/L	APHA 3120 B	5	2
3.	Arsenic (as As), µg/L	APHA 3120 B	10	Less than 4 ⁽²⁾
4.	Boron (as B), µg/L	APHA 3120 B	1000	Less than 5 ⁽²⁾
5.	Cadmium (as Cd), µg/L	APHA 3120 B	5	Less than 1 ⁽²⁾
6.	Chromium (as Cr), µg/L	APHA 3120 B	50	Less than 1 ⁽²⁾
7.	Iron (as Fe), µg/L	APHA 3120 B	200	Less than 0.4 ⁽²⁾
8.	Lead (as Pb), µg/L	APHA 3120 B	10	Less than 6 ⁽²⁾
9.	Manganese (as Mn), µg/L	APHA 3120 B	50	Less than 0.1 ⁽²⁾
10.	Mercury (as Hg), µg/L	CTS/TP/AE/ In-house 002	1	Less than 1 ⁽²⁾
11.	Nickel (as Ni), µg/L	APHA 3120 B	20	Less than 1 ⁽²⁾
12.	Selenium (as Se), µg/L	APHA 3120 B	10	Less than 2 ⁽²⁾

Note : ⁽¹⁾ Since result of test after the first 24 hours extraction comply with the requirements, this first extract is defined as the final extract.

⁽²⁾ Limit of detection

Comment: The sample tested complied with all the requirements of BS 6920-1: 2014: Clause 8

