

Test Report Page: 1 of 3 No.: CT/2017/61188A Date: 2017/08/03

BIOPLASTICOS Y POLIMEROS BIOFASE S.A.P.I DE C.V MADERO ORIENTE 6500, CIUDAD INDUSTRIAL, MORELIA MICHOACAN MEXICO, CP:58200

The following sample(s) was/were submitted and identified by/on behalf of the applicant as:

BIOPLASTICOS Y POLIMEROS BIOFASE S.A.P.I DE C.V Sample Submitted By

Sample Description **BIOBASED CUTLERY**

Sample Material **CUTLERY** Sample Receiving Date 2017/06/29

Testing Period 2017/06/29 TO 2017/07/13

: As specified by client, with reference to U.S. Food and Drug Administration (FDA) **Test Requested**

21 CFR 175.300 (Condition of use E) to conduct test. Please refer to result table

for testing item(s).

Test Result(s) Please refer to next page(s).



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BIOPLASTICOS Y POLIMEROS BIOFASE S.A.P.I DE C.V



MADERO ORIENTE 6500, CIUDAD INDUSTRIAL, MORELIA MICHOACAN MEXICO, CP:58200

Test Result(s)

PART NAME No.1 WHITE PLASTIC SPOON

PASS

Test Item (s)	Unit	Method	MDL	Result	Limit
				No.1	
Extractives residue (n-Heptane, 70°F, 30 min)	mg/in²	With reference to US FDA 21 CFR 175.300 condition E (2015).	0.1	n.d.	0.5
Extractives residue (D.I. Water, 120°F, 24 hr)	mg/in²	With reference to US FDA 21 CFR 175.300 condition E (2015).	0.1	0.142	0.5

Note:

- 1. 0.1wt% = 1000ppm; mg/kg = ppm
- 2. MDL = Method Detection Limit
- 3. n.d. = Not Detected
- 4. This report supersedes the previous document bearing the test report number CT/2017/61188.

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* The tested sample / part is marked by an arrow if it's shown on the photo. *

CT/2017/61188



** End of Report **

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