

Test Report No. SHAHG1724426601 Date: 14 Nov 2017 Page 1 of 4

IBOYA PACKAGING CO., LTD

NO.225, SUHONG ROAD, SUZHOU INDUSTRIAL PARK, SUZHOU, JIANGSU PROVINCE, CHINA,215021

The following sample(s) was/were submitted and identified on behalf of the clients as: 38-400 PLASTIC CAP,

WHITE PE LINER

SGS Job No.: SHHL1711062665OT - SH

Supplier: IBOYA PACKAGING CO., LTD

Country of Origin: CHINA

Date of Sample Received: 07 Nov 2017

Testing Period: 07 Nov 2017 - 14 Nov 2017

Test Requested : Selected test(s) as requested by client.

Test Method : Please refer to next page(s).

Test Results : Please refer to next page(s).

Result Summary:

Test Requested	Conclusion
FDA 21 CFR 177.1520- Extractable fraction	PASS
FDA 21 CFR 177.1520- Soluble fraction	PASS
FDA 21 CFR 177.1520- Density at 23°C	PASS
FDA 21 CFR 177.1520- Melting point	PASS

Signed for and on behalf of

Gerener Wanf

SGS-CSTC Standards Technical Services (Shanghai) Co., Ltd.

Serena Wang

Approved Signatory



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Test Results:

Test Part Description:

Specimen No. SGS Sample ID Description Material (claimed by the client)

SN1 SHA17-244266.001 Black plastic lid PP

SN2 SHA17-244266.002 White plastic washer PE

FDA 21 CFR 177.1520- Extractable fraction

Test Method: For sample 001: With reference to FDA 21 CFR 177.1520(d) 3 (i).

For sample 002: With reference to FDA 21 CFR 177.1520(d)(3)(ii).

Simulant Used	<u>Time</u>	<u>Temperature</u>	Max. Permissible	Result of 001	Conclusion
			<u>Limit</u>		
n-hexane	2hr(s)	Reflux temperature	6.4% (w/w)	0.7% (w/w)	PASS
Simulant Used	<u>Time</u>	<u>Temperature</u>	Max. Permissible <u>Limit</u>	Result of 002	Conclusion
n-hexane	2hr(s)	50°C	5.5% (w/w)	<0.5% (w/w)	PASS

FDA 21 CFR 177.1520- Soluble fraction

Test Method: For sample 001: With reference to FDA 21 CFR 177.1520 (d) 4 (i).

For sample 002: With reference to FDA 21 CFR 177.1520(d)(4)(ii).

Simulant Used	<u>Time</u>	<u>Temperature</u>	Max. Permissible	Result of 001	Conclusion
			<u>Limit</u>		
Xylene	-	25°C	9.8% (w/w)	3.4% (w/w)	PASS
Simulant Used	<u>Time</u>	<u>Temperature</u>	Max. Permissible <u>Limit</u>	Result of 002	Conclusion
Xvlene	_	25°C	11.3% (w/w)	4.0% (w/w)	PASS

FDA 21 CFR 177.1520- Density at 23°C

Test Method: With reference to FDA 21 CFR 177.1520 d (1).



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 Test Item(s)
 Limit
 001

 Density at 23°C, g/ cm³
 0.880-0.913
 0.900

 Conclusion
 PASS

FDA 21 CFR 177.1520- Melting point

Test Method: With reference to FDA 21 CFR 177.1520 (d) (2).

Test Item(s)Limit001Melting point, °C*160-180165ConclusionPASS

Notes:

(1) *The test was subcontracted to SGS Ningbo chemical lab.



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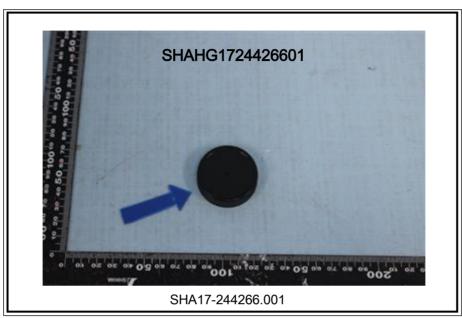
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Sample photo:





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*** End of Report ***



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