



# Certificate Of FDA Compliance

201601 -201701

This is certified that:

At The Address Stated Below Has Completed U.S. FOOD And DRUG  
ADMINISTRATION Food Facility Registration And Test Through MANTONG  
(www.fdacn.org)

**JINGJIANG OHJI RUBBER CO.,LTD**

**BAWEI TOWN,JINGJIANG CITY,JIANGSU PROV,CHINA**

**Product Description:**

**B-5BS**

**Report No.**

**J-00205498**

**Test Method:**

**FDA Food Contact Article tests in  
accordance with 21 CFR 177.2600**

**Test Performing Date:**

**17-Dec-2015 to 23-Dec-2015**

**Test Laboratory**

**NSF USA**

**Registration Number**

**19526685684**



*Jacky M. Chuang*

Executive Director

Date: 01-05-2016

**MTG STANDARDS TESTING & CERTIFICATION CENTER** [www.fdacn.org](http://www.fdacn.org)

*This certification affirms that the above device and company was registered with U.S. Food and Drug Administration pursuant to section 305 of the United States Public Health and Bioterrorism Preparedness and Response Act of 2002, P.L. 107-188, on the date stated above, and makes no other representations or warranties, nor does this certificate make any representations or warranties to any person or entity other than the named certificate holder, for whose sole benefit it is issued. MTG CO., Inc. assumes no liability to any person or entity in connection with the foregoing. MTG is a private registration agent not affiliated with the U.S. Food and Drug Administration.*

## NSF International

NSF International.  
789 Dixboro Road, Ann Arbor, Michigan  
48105-9723 USA  
1-800-NSF-MARK 734-769-8010  
www.nsf.org

# TEST REPORT

**Send To: C0096491**

**Facility: C0096493**

JINGJIANG OHJI RUBBER CO.,LTD  
BAWEI TOWN,JINGJIANG CITY,  
JIANGSU PROV,CHINA

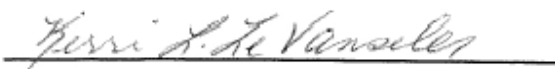
JINGJIANG OHJI RUBBER CO.,LTD  
BAWEI TOWN,JINGJIANG CITY,  
JIANGSU PROV,CHINA

Result	Complete	Report Date	23-Dec-2015
Customer Name	JINGJIANG OHJI RUBBER CO.,LTD		
Description	B-5BS		
Test Type	Test Only		
Job Number	J-00205498		
Sample Reception Date	17-Dec-2015		
Testing Completion Date	23-Dec-2015		

### Summary of Results

Testing Parameters and Standards	Result
Determination of extractive residue according to US FDA 21 CFR 177.2600	Complete

**Report Authorization**



Kerri Levanseler - Director, Chemistry Laboratory

**Date** 23-Dec-2015

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1. **Determination of extractives residue according to US FDA 21 CFR 177.2600**

1.1 Testing procedure:

Extract samples with the extractive solvents at the indicated extractive conditions (water at reflux temperature for initial 7h and later 2h, n-hexane at reflux temperature for initial 7h and later 2h), weigh the residue and calculate total extractives in mg per square inch of the sample extracted.

1.2 Testing result:

Sample No.	Testing Parameter	Unit	Result	Acceptance	Conclusion	
S-0001217120	Extractives residue ( Water )	Initial 7h	mg/in <sup>2</sup>	ND(1.0)	20	Pass
		Later 2h	mg/in <sup>2</sup>	ND(0.1)	1	Pass
	Extractives residue ( n-hexane )	Initial 7h	mg/in <sup>2</sup>	21	175	Pass
		Later 2h	mg/in <sup>2</sup>	0.7	4	Pass

Remark

- 1) ND=Not Detected, less than reporting limit.
- 2) Client claim the material of sample is Rubber.
- 3) End-product testing was performed to 21 CFR 177.2600 per Client request and that the acceptable results are based on the assumption that all material components have been confirmed by the formulator as compliant to 21 CFR and applicable FDA regulations for the intended end use.
- 4) Shanghai Laboratory was authorized by Ann Arbor to conduct the testing.

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**Picture of Sample**



B-5BS

**End of Report**