# Certificate Of Fda Registration

#MTGCHN09032502FDA#

## This is certified that:

**At The Address Stated Below Has Completed U.S. FOOD And DRUG ADMINISTRATION Food Facility Registration And Test Through MANTONG.**

**Meder-GHL Electronic Co., Ltd. Shanghai**

Add:F/2 #7 Building Zhangmu Industrial Zone #128
Yaobei Road Jiuting,Shanghai

| | |
|---|---|
| FDA Facility Number: | 16862986396 |
| Product Description: | PP Floater (KSS/MS– PP series) |
| Test Requested: | Tests for compliance with U.S. F.D.A. C.F.R. 21 Part 177.1520 (for Polypropylene homo-polymer resin in contact with food |
| Test Method: | With reference to U.S. F.D.A. C.F.R. 21 Part 177.1520. |
| Test Performing Date: | 2009-02-19 - 2009-02-27 |
| Report No. : | AL006132-001 |

MTG FDA REGISTER CENTER

Jacky M. Chuang

Executive Director

Date: 03/15/2009

MTG FDA REGISTER CENTER CORPORATE SEAL CALIFORNIA

This certification affirm 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 2002, P.L. 107-188, on the date stated above, and makes no other representations or warranties. It did on this certificate make any representations or warranties to any person or not us, other than the named certificate holder for which 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.