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Client BOOSTER(DONGGUAN)PLASTIC PRODUCTS CO.,LTD

**Client Address** NO.497 Tangxia North Road, TangXia Town, Dongguan City, Guangdong, China

The following sample(s) and sample information was/were submitted and confirmed by the client:

Sample Name Loop pin/Tag pin

**Specification** /

Material PP

\*\*\*\*\*\*

**Date of Sample** 

2019-11-13

Received

**Test Period** 2019-11-13~2020-01-21

Evaluation of potential biodegradation and disintegration of materials under the action of

Test Item(s)

specific microorganisms (soil landfill test)

Test Result(s) Please see the next page(s)

> Compiled by: Hardong Pan.

Inspected by:

Approved by:

**Issued Date:** 

Shanghai Micro-spectial Chemical Analysis and Test technology Co., Ltd.



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### 1. Test Item:

Test Standard: GB/T 19275-2003

Lab Environment: The experiment was conducted in a biological incubator at  $29\,^\circ\!\text{C} \pm 1\,^\circ\!\text{C}$  with relative

humidity of 95% ±2% for 28 days

Test Result(s):

The sample name	Test Item	Test Result	
Loop pin/Tag pin	Evaluation of potential biodegradation and disintegration of materials under the action of specific microorganisms (soil landfill test)	Change in mass	+0.01%



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### 2. Detection process

#### 2.1 Test conditions

- **2.1.1**Sample size: square with side length of 50mm×50mm;
- 2.1.2Properties of landfill soil: the water retention capacity is 78.9%, the water content is 60.9%, the pH value is
- 4.72, and the biological activity meets the requirements.
- **2.1.3** Sterilization solution: o-phenyl phenol;
- **2.1.4** Test methods: methods C soil landfill test (evaluation of quality change).

#### 2.2 Test group

- 2.2.1 Prepare 3 groups of samples, 6 samples for each group, a total of 18 samples;
- 2.2.1.1 Group 0: control samples, stored under standard temperature and humidity conditions;
- 2.2.1.2GroupI: samples inoculated with microorganisms and cultured (directly buried in soil);
- **2.2.1.3**Group S: sterilized samples were cultured under the same conditions as group I.

### 2.2.2 Biological activity of soil.

The bleached but untreated cotton fabric  $(250g/m^2)$  strip  $(2.5cm \times 10cm)$  was buried in the soil at the same time with the sample and cultured for 7 days. After the end of the cultivation period, the stretch length of the cotton strip should not be more than 25%, and the biological activity of the soil should meet the requirements.

**2.2.3**Fill 1 L jar with soil with water retention capacity with humidity of  $(60\% \pm 5\%)$ .

### 2.2.4Fill the samples

With a spoon and tweezers, the samples were buried in two jars, each containing a comparison strip of cotton to test the soil's activity (after the culture period, the stretch length of the strip should not be greater than 25%, and the soil's biological activity should meet the requirements). Each jar was filled with 3 samples vertically and arranged in parallel. A total of 6 samples of the two jars were used as group I samples. To get the oxygen flowing, a wire of about 1mm is wound between the jar and the cap. The soil in the jar shall not be compressed when filling the sample, and the soil covering the sample shall not be more than 12.5 cm thick.

### 2.2.5 Test inspection

The 1L sealed storage jar containing soil and cotton strips was placed in the autoclave sterilizer and sterilized continuously for 30min 3d at  $120^{\circ}$ C and  $2\times105$  Pa. The samples were first immersed in o-phenylphenol solution, then placed in two jars, and 3mL of o-phenylphenol solution was poured onto the soil. Each jar was filled with 3 samples, which were arranged in parallel with each other. The two jars contained 6 samples, which were used as group S samples. The sealing methods of jar caps and soil filling methods of group I and group S were consistent.

### 2.2.6 The process of cultivation

The prepared two jars in group I and group S were placed in the incubator and cultured for 28 days in the biological incubator with the temperature of  $29^{\circ}\text{C} \pm 1^{\circ}\text{C}$  and the relative humidity of  $95\% \pm 2\%$ .

#### 2.2.7 The measurement process

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### 2.2.7.1 Cleaning steps

After the experiment, the samples of group I, group S and group 0 were immersed in 70% ethanol solution for about 5min, then rinsed under running water, wiped with filter paper, and placed in a petri dish with no cover. Then the samples were dried to constant weight in a dryer together with the petri dish.

### 2.2.7.2 Quality of change

Cleaning: Dip the test specimens for 5 min into an 70% ethanol-water mixture, rinse under running water, wipe with filter paper place it into a clean petri dish and allow to dry overnight at room temperature. And then put it in a desiccator until constant mass is attained.

Changes in mass: For the determination of the change in mass, place the cleaned specimens in a desiccator and weight them regularly to the nearest 0.1mg until constant mass is attained. Record the final masses. For each specimen, the difference in mass  $\Delta$  m before and after exposure. The average percentage change in mass using the formula:

$$(\overline{\Delta m I} - \overline{\Delta m S}) / m e \times 100$$

Where  $\Delta$  m I and  $\Delta$  m S is the arithmetic mean of the changes in mass of test series I and S. m e is the mean of the original specimen masses.

## 2.3 The results

## 2.3.1 The resultsof quality change

The results of quality changes of samples in each group are shown in Table 1. According to the results in Table 1, the sample quality change is +0.01%.

## Table1Theresultsof quality change

Group			Gro	oupI		GroupS						Group0				
Project	The initial qualitym <sub>I</sub> (mg)		The final qualitym <sub>I</sub> '		The quality change after $test \Delta m_I$ (mg)		e initial alitym <sub>s</sub>	qual	e final itym <sub>S</sub> ' mg)	The quality change after test∆m <sub>S</sub> (mg)	The initial qualitym <sub>0</sub> (mg)		The final qualitym <sub>0</sub> ' (mg)		The quality change after test∆m <sub>0</sub> (mg)	
The mass	$m_1$	627.37	$m_1{'}$	628.31	0.94	$m_1$	281.03	$m_1{'}$	281.37	0.34	$m_1$	281.79	$m_1{'}$	281.96	0.17	

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after	$m_2$	637.72	$m_2'$	638.23	0.51	$m_2$	638.04	$m_2'$	639.06	1.02	$m_2$	633.75	$m_2'$	634.19	0.44
constant	$m_3$	276.82	$m_3$	277.58	0.76	$m_3$	278.62	$m_3$	279.55	0.93	$m_3$	285.92	$m_3$	286.13	0.21
weight	$m_4$	630.47	$m_4{}'$	631.02	0.55	m <sub>4</sub>	635.73	$m_4{}'$	636.69	0.96	m <sub>4</sub>	266.71	$m_4$	266.88	0.17
	$m_5$	286.91	$m_5$	287.77	0.86	$m_5$	277.22	$m_5$	277.99	0.77	$m_5$	630.50	$m_5$	631.25	0.75
	$m_6$	280.64	$m_6$	281.93	1.29	$m_6$	627.51	$m_6$	628.17	0.66	$m_6$	633.27	$m_6$	633.61	0.34
Average result	`m <sub>I</sub>	456.66	`m <sub>I</sub> '	457.47	0.82	`m <sub>S</sub>	456.36	`m <sub>S</sub> ′	457.14	0.78	`m <sub>0</sub>	455.32	$m_0'$	455.67	0.35
$m_{\rm e}$ (The mean mass of the original sample) (mg)						456.11									
The average percentage change in mass (%)							0.01								

### Remark:

<sup>1.</sup> In the table, mis the quality before the test, m is the quality after the test, and  $\Delta m$  is the quality difference before and after the test.  $m_e$  is the mean value of original sample quality.

<sup>2.</sup> Please refer to the attached drawings for sample changes before and after the test.



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## 3. Representative attachments: pictures of samples before and after the test

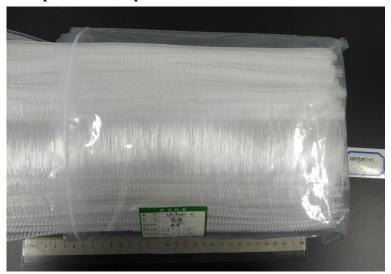


Fig. 1: Specimens before test

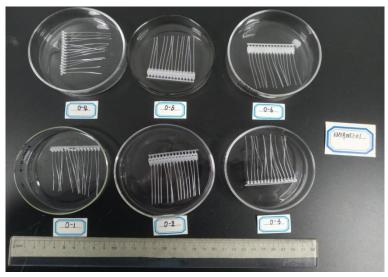


Fig. 2: Specimens of test series 0 before test



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Fig. 3: Specimens of test series S before test



Fig. 4: Specimens of test series I before test



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Fig. 5: Specimens of test series S after test



Fig. 6: Specimens of test series I after test



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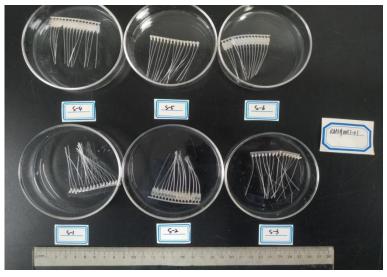


Fig. 7: Specimens of washed test series S after test



Fig. 8: Specimens of washed test series I after test



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## 4. Sample photos:



\*\*\* End of the Report\*\*\*

#### **DECLARE**

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