

Report No. SCL03I000625011

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ApplicantZHONGSHAN FUZHOU ADHESIVE-PRODUCT CO.,LTDAddressTORCH HIGH-TECH DEVELOPMENT ZONE,ZHONGSHAN

CITY, GUANGDONGPROVINCE CHINA

The following sample(s) and sample information was/were submitted and identified by/on the behalf of the client

Sample Name PET镀铝膜 PET Aluminum film

Client Reference TL-S 25#/TL-S 50#/TL-S 75#/TL-S 100#/TL-SM 25#/TL-SM 50#/
Information TL-SM 75#/TL-SM 100#/TL-SM 125#/TL-SM 150#/TL-SM 200#/
TL-SMB 25#/TL-SMB 50#/TL-SMB 75#/TL-SMB 100#/TL-SH 25#/

TL-SMB 25#/TL-SMB 50#/TL-SMB 75#/TL-SMB 100#/TL-SH 25#/ TL-SH 50#//TL-SH 100#/TL-G 25#/TL-G 50#/TL-G 100#/TL-GM 25#/ TL-GM 50#/TL-GM 100#/TL-GMB 25#/TL-GMB 50#/TL-GMB 100#/

TL-GH 25#/TL-GH 50#//TL-GH 100#

Sample Received Date Jan. 30, 2016

Testing Period Jan. 30, 2016 to Feb. 5, 2016

Test Requested As specified by client, to test Polycyclic Aromatic Hydrocarbons(PAHs) in the

submitted sample(s).

Test Method

Test Item(s)	Test Method	Measured Equipment(s)
Polycyclic Aromatic Hydrocarbons(PAHs)	AfPS GS 2014:01 PAK	GC-MS

Test Result(s)

Please refer to the following page(s).

Conclusion

Tested Item	Standard	Result
Polycyclic Aromatic Hydrocarbons (PAHs)	AfPS GS 2014:01 PAK, Category 3	Pass

Too Chen

Ginghong Wan

Qinghong Wan

Technical Manager

Reviewed by Wenjun Wang

Date Feb. 6, 2016

No. R131024944

Centre Testing International Group Co.,Ltd. Shunde Branching Building, Section 2, No.8, East of Rongqi Avenue, Ronggui, Shunde District, Foshan, Guangdong, China

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Limits for PAHs content (mg/kg) for material of (grip) surfaces, which are to be categorized on account of the results of the risk analysis.

	Category 1	Categor	y 2	Categor	ry 3
Parameters	Materials intended to be put in the mouth or materials of toys with foreseeable long-term skin contact (longer	Materials not covered by category 1, with foreseeable skin contact for longer than 30 seconds (long-termskin contact) or repeated # short-term skin contact		Materials not covered by category 1 or 2 with foreseeable skin contact up to 30 seconds (short termskin contact)	
	than 30 seconds)	Toys covered by Directive 2009/48/EC	Other products	Toys covered by Directive 2009/48/EC	Other products
Benzo[a]pyrene	<0.2	<0.2	<0.5	<0.5	<1
Benzo[e]pyrene	<0.2	< 0.2	< 0.5	< 0.5	<1
Benzo[a]anthracene	<0.2	<0.2	<0.5	< 0.5	<1
Benzo[b]fluoranthene	<0.2	< 0.2	<0.5	< 0.5	<1
Benzo[j]fluoranthene	<0.2	< 0.2	< 0.5	<0.5	<1
Benzo[k]fluoranthene	<0.2	<0.2	<0.5	< 0.5	<1
Chrysene	<0.2	<0.2	<0.5	< 0.5	<1
Dibenz[a,h]anthracene	<0.2	<0.2	<0.5	<0.5	<1
Benzo[g,h,i]perylene	<0.2	<0.2	<0.5	< 0.5	<1
Indenol[1,2,3-cd]pyrene	<0.2	<0.2	< 0.5	< 0.5	<1
Acenaphthylene, Acenaphthene, Fluorene, Phenanthrene, Anthracene, Fluoranthene Pyrene	<1 Sum	<5 Sum	<10 Sum	<20 Sum	<50 Sum
Naphthalene	<1	<2		<10	
Sum 18 PAHs	<1	<5	<10	<20	<50

Formulation "of repeated short-term skin contact" REACH Annex XVII No. 50 supplement (REGULATION (EU) No.1272/2013)





















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Test Result(s)

Tested Item(s)	Result	MDL
Polycyclic Aromatic Hydrocarbons (PAHs)	(45)
Naphthalene	N.D.	0.2 mg/kg
Acenaphthylene	N.D.	0.2 mg/kg
Acenaphthene	N.D.	0.2 mg/kg
Fluorene	N.D.	0.2 mg/kg
Phenanthrene	N.D.	0.2 mg/kg
Anthracene	N.D.	0.2 mg/kg
Fluoranthene	N.D.	0.2 mg/kg
Pyrene	N.D.	0.2 mg/kg
Benzo[a]anthracene	N.D.	0.2 mg/kg
Chrysene	N.D.	0.2 mg/kg
Benzo[b]fluoranthene	N.D.	0.2 mg/kg
Benzo[k]fluoranthene	N.D.	0.2 mg/kg
Benzo[a]pyrene	N.D.	0.2 mg/kg
Indeno[1,2,3-cd]pyrene	N.D.	0.2 mg/kg
Dibenzo[a,h]anthracene	N.D.	0.2 mg/kg
Benzo[g,h,i]perylene	N.D.	0.2 mg/kg
Benzo[j]fluoranthene	N.D.	0.2 mg/kg
Benzo[e]pyrene	N.D.	0.2 mg/kg
Sum (Acenaphthylene, Acenaphthene, Fluorene, Phenanthrene, Anthracene, Fluoranthene, Pyrene)	N.D.	/
Sum 18 PAHs	N.D.	

Tested Sample/Part Description

Silvery plastic film

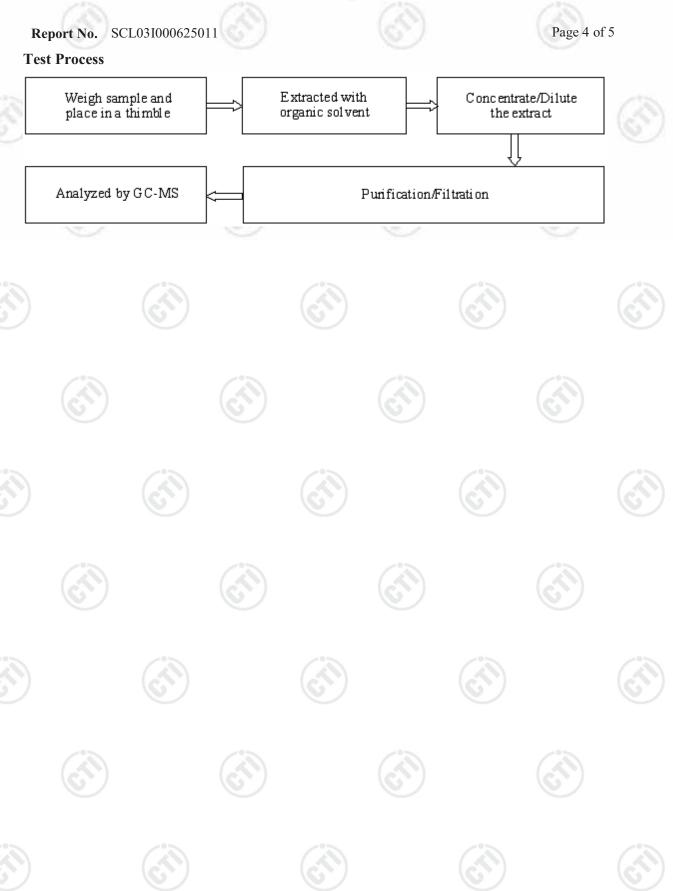
Remark: -MDL = Method Detection Limit

-N.D. = Not Detected (<MDL)

-mg/kg = ppm = parts per million

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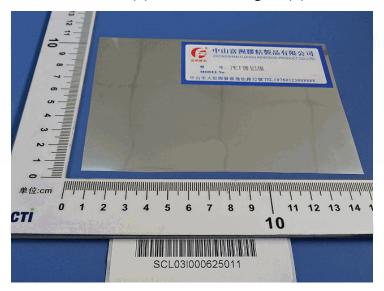




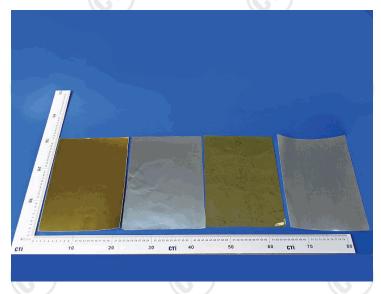
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Photo(s) of the sample(s)



Photo(s) for reference



At the request of client, the product(s) in this report is/are provided by applicant (non-tested sample(s)).

*** End of report ***

The test report is effective only with both signature and specialized stamp. The result(s) shown in this report refer only to the sample(s) tested. Without written approval of CTI, this report can't be reproduced except in full.

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