



# TEST REPORT

Test Report # 18H-005439 Date of Report Issue: August 8, 2018  
 Date of Sample Received: July 23, 2018 Pages: Page 1 of 8

## CLIENT INFORMATION:

Company: Calibre International, LLC  
 Recipient: Lisa Arismendez  
 Recipient Email: LArismendez@highcaliberline.com



## SAMPLE INFORMATION:

Description:	1oz. Hand Sanitizer Gel w/carabiner		
Assortment:	-	Purchase Order Number:	-
SKU/style No.:	H304	Toy Co./Agency:	-
Factory/Supplier/Vendor:	-	Country of Origin:	China
Country of Distribution:	United States	Labeled Age Grade:	-
Quantity Submitted:	3 pcs	Recommended Age Grade:	-
Testing Period:	07/25/2018 – 08/06/2018	Tested Age Grade:	-

## OVERALL RESULT:

**PASS**

Refer to page 2 for test result summary and appropriate notes.

ANSECO GROUP (HK) LIMITED

Loska Yeung Lok Ka  
 Assistant Manager, Chemical Laboratory

ANSECO GROUP (HK) LIMITED • 3/F Liven House • No. 61 – 63 King Yip Street • Kwun Tong • Kowloon • Hong Kong • Tel: (852)3185 8000

The above test(s) is/are accredited under the laboratory's ISO/IEC 17025 accreditation issued by the ANSI-ASQ National Accreditation Board (ANAB) according to certificate and scope of accreditation (Certificate # AT-1500.) Test(s) marked with '#' is/are not covered under the scope of accreditation.

The test result(s) and conclusion(s) in this report relate to the sample(s) tested as described herein.

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**TEST RESULTS SUMMARY:**

At the request of the client, the following tests were conducted:

CONCLUSION	TEST(S) CONDUCTED
PASS	California Proposition 65, Total Lead in Substrate Materials
PASS	California Proposition 65, Phthalates (DBP, BBP, DEHP, DINP, DIDP, DnHP)
Refer to detailed result	FDA 21 CFR Part 201.62 (q), Net Quantity of Contents <sup>#</sup>
Refer to detailed result	Client's Requirement, Alcohol Content <sup>#</sup>

**DETAILED RESULTS:****California Proposition 65, Total Lead in Substrate Materials**

Test Method: CPSC-CH-E1001-08.3 (Metal), CPSC-CH-E1002-08.3 (Non-Metal)

Analytical Method: Inductively Coupled Plasma-Optical Emission Spectrometry

Specimen No.	1+2	3	4	5	6	Total Limit (ppm)
Test Item	Result (ppm)	Result (ppm)	Result (ppm)	Result (ppm)	Result (ppm)	
Total Lead (Pb)	ND	ND	ND	ND	ND	<b>100</b>
<b>Conclusion</b>	PASS	PASS	PASS	PASS	PASS	

Specimen No.	7	8	---	---	---	Total Limit (ppm)
Test Item	Result (ppm)	Result (ppm)	Result (ppm)	Result (ppm)	Result (ppm)	
Total Lead (Pb)	ND	ND	---	---	---	<b>100</b>
<b>Conclusion</b>	PASS	PASS	---	---	---	

**Note:**

ppm (Parts per million) = mg/kg (Milligrams per kilogram)

LT = Less than

ND = Not detected (Reporting Limit = 20 ppm)

Composite results are based on specimen of least mass resulting in highest potential concentration.

**Remark:**

The specification is quoted from client's requirement.

**DETAILED RESULTS:****California Proposition 65, Phthalates (DBP, BBP, DEHP, DINP, DIDP, DnHP)**

Test Method: CPSC-CH-C1001-09.3

Analytical Method: Gas Chromatography with Mass Spectrometry

Specimen No.		1+2	3	4	---	Limit (ppm)
Test Item	CAS No.	Result (ppm)	Result (ppm)	Result (ppm)	Result (ppm)	
Dibutyl phthalate (DBP)	84-74-2	ND	ND	ND	---	<b>1000</b>
Benzyl butyl phthalate (BBP)	85-68-7	ND	ND	ND	---	<b>1000</b>
Di-(2-ethylhexyl) phthalate (DEHP)	117-81-7	ND	ND	ND	---	<b>1000</b>
Diisononyl phthalate (DINP)	28553-12-0 68515-48-0	ND	ND	ND	---	<b>1000</b>
Diisodecyl phthalate (DIDP)	26761-40-0 68515-49-1	ND	ND	ND	---	<b>1000</b>
Di-n-hexyl phthalate (DnHP)	84-75-3	ND	ND	ND	---	<b>1000</b>
<b>Conclusion</b>		PASS	PASS	PASS	---	

**Note:**

ppm (Parts per million) = mg/kg (Milligrams per kilogram) = 0.0001 % m/m (Percent by mass)

LT = Less than

ND = Not detected (Reporting Limit = 120 ppm)

Composite results are based on specimen of least mass resulting in highest potential concentration.

**Remark:**

The specification is quoted from client's requirement.



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**DETAILED RESULTS:****FDA 21 CFR Part 201.62 (q), Net Quantity of Contents<sup>#</sup>**

Specimen	Result	Labelled Content	Conclusion
8	0.6591 fl oz	1.051 fl oz	Information only

NA = Not applicable

fl oz = Fluid ounces

## Remark:

\*Client's specification. From 21 CFR Part 201.62(q), the declaration of net quantity of contents shall express an accurate statement of the quantity of contents of the package. Reasonable variations caused by loss or gain of moisture during the course of good distribution practice or by unavoidable deviations in good manufacturing practice will be recognized. Variations from stated quantity of contents shall not be unreasonably large.



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**DETAILED RESULTS:****Client's Requirement, Alcohol Content**

Test Method: In-House Method#  
Analytical Method: Gas Chromatography

Specimen No.	Result	Conclusion
8	49.3%	Information only

*Note:*

% = Percentage by weight

**SPECIMEN DESCRIPTION:**

Specimen No.	Specimen Description	Location
1	Translucent plastic	Lid
2	Transparent plastic	Body
3	Multicolor printed white/ grey PVC with adhesive	Plastic sticker
4	Blue printed clear plastic with adhesive	Plastic sticker
5	Silvery metal	Carabiner
6	Matt silvery metal	Opener of carabiner
7	Off silvery metal	Rivet
8	Transparent gel	Gel



**SAMPLE PHOTO:**



-End Report-