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Applicant:	Vandy Vape			
Address:	415, huafeng Internet	+ creative park, no.107	republic industrial r	oad,
	xixiang, baoan distric	t, shenzhen		
The following sample w	as submitted and iden	tified by/on behalf of th	e client as:	
Sample Name:	Bonza Kit			
Model No.:	Bonza Kit			
Tank:	Juice Capacity 2ml,	SS304		
Power level in testing:	Voltage/Wattage of t	ested sample is un-adjust	table	
Adjustable air inlet or not:	Yes			
Trade Mark:	Vandy Vape			
Sample Received Date:	2018.06.25			
Testing Period:	2018.06.25—2018.0	7.03		
Test Method:	Please refer to the fo	ollowing page(s).		
Test Result(s):	Please refer to the fo	ollowing page(s).		

Tes	at Items	Test Requested
1	Carbonyl Compounds: Formaldehyde, Acetaldehyde, Acrolein, Crotonaldehyde	Emission testing according to
2	Metals: Aluminum, Chromium, Iron, Nickel, Tin, Lead, Cadmium, Arsenic, Antimony	Article 20 of
		Tobacco Product
3	Nicotine consistency	Directive
		(2014/40/EU)

Checked by Signed for and on behalf of TCT Yir Noel Noel Yin Kim Zhang **Technical Manager**

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Test Results:

Test Condition for test items except Nicotine consistency test:

With reference to the CORESTA RECOMMENDED METHOD Nº 81 method parameter and Afnor standardization XP D90-300-3, a smoke machine was used to collect the vapor.

	Puff Duration	3.0s±0.1s	
	Puff Volume	55mL±0.3mL	
	Puff Frequency	30s±0.5s	
<u> </u>	Puff of Each Group	20	T _N C)
	Group Interval Time	300s±120s	
	Maximum Flow	18.5mL/s±1.0mL/s	
	Pressure Drop	< 50hPa	
	Group	5	
	Total Number of Puff	100	
<u>(</u> ()	Total Duration of Vaporization	300s	10

The temperature and relative humidity of the test atmosphere during machine preparation and testing were kept within the following limits: temperature $\pm 2^{\circ}$ C, relative humidity $\pm 5\%$





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1. Carbonyl Compounds Content(s)

Method: The volatile aldehydes are extracted from the aerosol by bubbling each puff through an impactor containing an acidified aqueous solution of 2,4-DNPH. The samples are analyzed by reverse phase high-performance liquid chromatography and determined using a UV detector.

Test Item	CAS No. Unit	Unit	MDI		Content(s)		
Test item	CAS NU.	Unit	MDL LOQ		No.1		
Formaldehyde	50-00-0	ug/100puffs	0.667	2	5.68		
Acetaldehyde	75-07-0	ug/100puffs	0.667	2	5.11		
Acrolein	107-02-8	ug/100puffs	0.667	2	ND		
Crotonaldehyde	4170-30-3	ug/100puffs	0.667	2	ND		

- Note: ug = Microgram
 - ND = Not Detected (lower than MDL)
 - MDL = Method Detection Limit
 - LOQ = Limit of Quantitation
 - E-Liquid Used: E-liquid B (AFNOR XP D90-300-3)



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2. Metals Content(s)

Method: The vapor was passed through a dry-ice cooled impinger containing glass packing beads and quartz wool. After smoking the impinger was extracted with 5% nitric acid and filtered through quartz wool. An aliquot of the resulting solution was submitted for analysis by ICP-OES.

Test Item CAS No		Linit	MDI		Content(s)		
rest tiem	CAS No.	Unit	MDL	LOQ	No.1		
Aluminum(Al)	7429-90-5	ug/100puffs	0.025	0.25	ND		
Chromium(Cr)	7440-47-3	ug/100puffs	0.005	0.05	ND VO		
Iron(Fe)	7439-89-6	ug/100puffs	0.005	0.05	ND		
Nickel(Ni)	7440-02-0	ug/100puffs	0.025	0.25	ND		
Tin(Sn)	7440-31-5	ug/100puffs	0.25	2.5	ND		
Lead(Pb)	7439-92-1	ug/100puffs	0.025	0.25	ND		
Cadmium(Cd)	7440-43-9	ug/100puffs	0.005	0.05	ND		
Arsenic(As)	7440-38-2	ug/100puffs	0.025	0.25	ND VO		
Antimony(Sb)	7440-36-0	ug/100puffs	0.025	0.25	ND		

Note: - ug = Microgram

- ND = Not Detected (lower than MDL)
- MDL = Method Detection Limit
- LOQ = Limit of Quantitation
- E-Liquid Used: E-liquid B (AFNOR XP D90-300-3)



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3. Nicotine Consistency Test

Test Condition: With reference to the CORESTA RECOMMENDED METHOD Nº 81 method parameter and Afnor standardization XP D90-300-3, a smoke machine was used to collect the vapor.

Puff Duration	3.0s±0.1s			
Puff Volume	55mL±0.3mL			
Puff of Each Group	20			
Maximum Flow	18.5mL/s±1.0mL/s			
Pressure Drop	< 50hPa			

The temperature and relative humidity of the test atmosphere during machine preparation and testing were kept within the following limits: temperature $\pm 2^{\circ}$, relative humidity $\pm 5\%$

Method: A reference liquid was prepared. A pharmaceutical nicotine inhaler was used as a comparator. Products were attached to a smoke machine, and the aerosol was collected in Cambridge filter pads. After trapping and solvent extraction, solution was analyzed by GC-MS and nicotine was dosed by comparing the areas obtained on the MS detector with those of standard solutions prepared in the laboratory under concentration conditions surrounding those of the samples.

Sample No.	Nicotine(CAS No.:54-11-5) Contents(mg/20Puffs)						Total
Sample No.	Group 1*	Group 2	Group 3*	Group 4	Group 5*	AVG	(mg/100puffs)
No.1	0.476	0.437	0.412	0.408	0.419	0.431	2.15
Deviation(%)	10.5	g -	4.2	-	2.6	-	
		1					

Note: - mg = milligram

- ND = Not Detected (lower than MDL)
- MDL = Method Detection Limit = 0.01mg/20Puffs
- LOQ = Limit of Quantitation = 0.1mg/20Puffs
- 1group = 20puffs
- * Values used for determination of consistency of nicotine emission
- E-Liquid Used: E-liquid A (AFNOR XP D90-300-3)
- Under the conditions of the test and with reference to AFNOR XP D90-300-3, the electronic cigarette delivers a dose of nicotine at consistent levels.



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