



Report Number: QIP-ASI179076

Audit Date : 23 Feb.,2017

This report is issued by Focus Technology Co., Ltd. (Made-in-China.com) and the supervising inspectorate (SGS-CSTC Standards Technical Services Co., Ltd.) to confirm that:

Company Name	: Wuhan Vanzpharm Inc. 武汉万知化工医药有限公司
Showroom	: http://vanzpharm.en.made-in-china.com
Address	 Room 2811, Jinliwu Mansion, Wuhan Economic & Technological Development Zone, Wuhan, Hubei, China Fanhu Industrial Park, Panwan, Jiayu County, Xianning, Hubei, China
Product	: Pharmaceutical Intermediate, API, Other Chemicals, Herbal Extract, Ginkgo Biloba Extract, Vitex Negundo Extract, Hypericum Perforatum Extract, Salvia Miltiorrhiza Extract, Hawthorn Extract, Forsythia Suspensa Extract, Motherwort Extract

has been on site audited for the Following Scope of Activity

- 1. General Information
- 2. Foreign Trade Capacity
- 3. Product Research & Development Capacity
- 4. Production Capacity & Quality Control
- ---the audit content of this part relates only to its associated enterprise
- 5. Working Environment
- 6. Photos

General Comments:



Wuhan Vanzpharm Inc. is a manufacturer company with 23 employees; it was established in 2011, located in Room 2811, Jinliwu Mansion, Wuhan Economic & Technological Development Zone, Wuhan, Hubei, China Fanhu Industrial Park, Panwan, Jiayu County, Xianning, Hubei, China. The lands occupy an area of 11790.44 square meters, and the annual production output was 420000 kg of pharmaceutical intermediate, 347 kg of API in last year. Wuhan Vanzpharm Inc. has its own brand. They have obtained 2 pcs of invention patent certificates for related products. Wuhan Vanzpharm Inc. has successful foreign trade experience in North America, South America, Europe, Southeast Asia/Mideast, East Asia & Australia.

Sign for and on behalf of SGS-CSTC Standards Technical Services Co., Ltd.



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SUPPLIER ASSESSMENT REPORT

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	Audited Company	Wuhan Vanzpharm Inc.				
	Audited Site:	Room 2811, Jinliwu Mansion, Wuhan Econon Wuhan, Hubei, China Fanhu Industrial Park, I China				
	Consigner of Assessment	Made-in-China.com				
	Audit Type	🛛 Initial Audit		III III B		
		Re-audit	胡北當色贝特美史 科技有限公司			
	Audit Date	23 Feb.,2017	Verify Report	www.sgs.com/ecv		
	Auditor	James Ai	Reviewed by	Simon Wang		
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	Part A: General Infor	mation				
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 Co.,Lid:
 新F.Haviling.jini.gavin





Part A: General Information

Section 1: Company Overview

1.1 Legal Validity				
Does the company have a valid business license?	🛛 Yes 🗌 No 📄 Others	Registration Number	91421221584883002B	
Year of established	22 Nov.,2011	Valid Date	21 Nov.,2021	
Registered address	Fanhu Industrial Park, Panwan, Jia	yu County, Xiann	ing, Hubei, China	
Actual address	Room 2811, Jinliwu Mansion, Wuh Wuhan, Hubei, China Fanhu Indus China			
Does the company in abnormal operation status list of industrial and commercial bureau?	No			
Registered capital	1000000 RMB			
Name of legal representative	Ms. Xiaolan Du			
Business scope	Research and Development, Whol pharmaceutical chemical products and explosive dangerous products and retails of chemical equipment; business of all kinds of goods a projects, needed approval and lice	(exclude the pois and pharmaceut Self-managemer and technology.(T	onous and harmful; inflammabl tical raw materials; Whole sale it and agency of import & expo The country has special limite	
1.2 Basic Information				
Contact person	Mr. Duke Du			
Phone number	0086-27-84492310	Fax number	0086-27-84492310	
URL/Web address	www.vanzpharm.com	1		
Company type		Company Corporation		
Type of ownership	Limited Company Public Company Foreign joint venture State-Owned Private Owner Wholly foreign-owned enterprises			
Associated company	 Audited company : Wuhan Vanz Associated company: Hubei Jiay The relationship of two companies	pharm Inc. vu Better Organic es: same shareholder	Co., Ltd.	
Products manufactured / sold scope	Pharmaceutical Intermediate, API, Extract, Vitex Negundo Extract, H Extract, Hawthorn Extract, Forsythi	hypericum Perfora	atum Extract, Salvia Miltiorrhiz	
1.3 Company Building Information) 1			
	estate certificate rvation Estimated on site			
The company area 4181.11 square m The land occupies 11790.44 square m The offices occupy 200 square meter The workshops occupy 3981.11 square	neters. s.			
いたいのでの した。 した。 した。 した。 した。 した。 した。 した。	erwise agreed in writing, this document is issued by th <u>sgs.com/en/Terms-and-Conditions.aspx</u> Attention is dra redin, Any holder of this document is advised that inform. readily accessible portions of the consignment and refil imits of Client's instructions, if any. The Company's sole transaction from exercising all their rights and obligati alsification of the content or appearance of this documen To check the authenticity of tasting /inspection reports Ch.Doccheck@sgs.com Din (Baixi) Robability Robability Robability (Nampular) Din (Baixi) Probability Robability (Nampular) Din (Baixi) Probability (Nampular) 	wn to the limitation of liabili ation contained hereon is s ects the Company's finding responsibility is to its Clie ions under the transaction it is unlawful and offenders rb ⁸ . certificate, please cor	ty, indemnification and jurisdiction issues olely limited to visual examination of the sat the time of its intervention only and than dhis document does not exonerate documents. Any unauthorized alteration, may be prosecuted to the fullest extent of stact us at telephone: (86-755) 8307 1443,	





Section 2: Human Resources

2.1 Company Chart					
2.2 Explanation of Co	C D E				
Code		Department		Number of employees	
A	General Manager			1	
В	Vice Manager			1	
С	Financial Dept.				
D	Sales Dept.			6	
E	Technology Dept.			2	
F	Admin & HR Dept.			1	
G	Production Dept.			11	
Н	Warehouse Dept.			1	
I	QC Dept.			1	
		Num	ber in total:	25	
2.3 Key Staff					
Title	Full name	Education	Working ex	xperience for this trade / total experience	
General Manager	Mr. Xiaolan Du	Doctor's Degree	15 / 15 years	s	

Remark:all above information based on staff lists, attendance records sheets and resume of key staff provided by company representative.



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Part B: Foreign Trade Capacity

Section 1: Export Overall Situation

1.1 Export Overall Situation	
Does the company have a valid Import and Export license?	🛛 Yes 🗌 No
The import and export enterprise code.	91421221584883002B
The number of foreign trading staff with relevant trading experience.	 □ within 1 year staff □ 6-10 years 2 staffs □ 0 years 2 staffs □ 0 years 1 staff □ 0 years 1 staff
The language freely used by foreign trade staff	🛛 English 🗌 others:
Annual revenue of previous year	Confidential
Annual export revenue of previous year	Confidential
Estimated export revenue for this year.	USD 1000000
Overseas agent / branch	🗌 Yes 🛛 No
Nearest port	Shanghai Port, Shenzhen Port
Acceptable quotation terms	🖾 FOB 🛛 CIF 🖾 CFR Other: EXW
Acceptable payment terms	 ☑ LC ☑ T/T ☑ D/P ☑ PayPal ☑ Western Union ☑ Small-amount payment
Average lead time (Peak Season)	□ within 15 workday ⊠ one month □ 2-3 months □ 4-6 months □ 6-12 months □ more time
Average lead time (Off Season)	☑ within 15 workday □ one month □ 2-3 months □ 4-6 months □ 6-12 months □ more time

Section 2: Export Business Capacity

2.1 Market Distribution (please list top three areas)				
Market	Main Product	Main client		
⊠ North America	Pharmaceutical Intermediate, API, Other Chemicals, Herbal Extract, Ginkgo Biloba Extract, Vitex Negundo Extract, Hypericum Perforatum Extract, Salvia Miltiorrhiza Extract, Hawthorn Extract, Forsythia Suspensa Extract, Motherwort Extract	Confidential		
South America	Pharmaceutical Intermediate, API, Other Chemicals, Herbal Extract, Ginkgo Biloba Extract, Vitex Negundo Extract, Hypericum Perforatum Extract, Salvia Miltiorrhiza Extract, Hawthorn Extract, Forsythia Suspensa Extract, Motherwort Extract	Confidential		
Europe	Pharmaceutical Intermediate, API, Other Chemicals, Herbal Extract, Ginkgo Biloba Extract, Vitex Negundo Extract, Hypericum Perforatum Extract, Salvia Miltiorrhiza Extract, Hawthorn Extract, Forsythia Suspensa Extract, Motherwort Extract	Confidential		



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Connecting Buyers with Chinese Suppliers

Southeast Asia/ Mideast	Pharmaceutical Intermediate, API, Other Chemicals, Herbal Extract, Ginkgo Biloba Extract, Vitex Negundo Extract, Hypericum Perforatum Extract, Salvia Miltiorrhiza Extract, Hawthorn Extract, Forsythia Suspensa Extract, Motherwort Extract	Confidential
Africa		
⊠ East Asia (Japan/ South Korea)	Pharmaceutical Intermediate, API, Other Chemicals, Herbal Extract, Ginkgo Biloba Extract, Vitex Negundo Extract, Hypericum Perforatum Extract, Salvia Miltiorrhiza Extract, Hawthorn Extract, Forsythia Suspensa Extract, Motherwort Extract	Confidential
⊠ Australia	Pharmaceutical Intermediate, API, Other Chemicals, Herbal Extract, Ginkgo Biloba Extract, Vitex Negundo Extract, Hypericum Perforatum Extract, Salvia Miltiorrhiza Extract, Hawthorn Extract, Forsythia Suspensa Extract, Motherwort Extract	Confidential
⊠ Domestic	Pharmaceutical Intermediate, API, Other Chemicals, Herbal Extract, Ginkgo Biloba Extract, Vitex Negundo Extract, Hypericum Perforatum Extract, Salvia Miltiorrhiza Extract, Hawthorn Extract, Forsythia Suspensa Extract, Motherwort Extract	Confidential
Others		



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Section 3: Supplier Management

3.1 Supplier Management						
ltem	Content	Observations /Comments				
1	Does the company establish and implement an effective suppliers' assessment procedure?	 Have the written procedures and followed records Have the written procedures but no records Have relevant records without written procedure No written procedures or followed records Other Remark: The auditor noted that there were some long-term suppliers to corporate. 				
2	Does the company have an updated list of approved suppliers?	 The approved suppliers list was updated in Have the written suppliers without approved signature or date. Provided some suppliers names No approved suppliers list Other Remark: The auditor noted that the company didn't provide approved suppliers due to commercial secret. 				

Section 4: After-sales Service Capacity

4.1 After-sales Service Capacity					
ltem	Content	Observations /Comments			
1	Is there a procedure to conduct random product inspection after final packaging in place?	 Have clear standards and written inspection records No written standards but had inspection reports Have the procedures but no inspection records It's not necessary to carry out the inspection Other Remark: The auditor noted that all final products were inspected before packaging on site, and also could accept to final random inspected before loading according to clients' requirement. 			
2	Is there a clear procedure for handling customer complaints?	 Has the clear procedure and followed records Has the procedure but no written records. No written procedures or records. Other Remark: The auditor noted that all customer complaints were followed by indicated sales with Email, Tel., Skype. 			
3	Can the finished/packaged product be traced by lot identification to the appropriate raw materials test reports?	 Have the procedures to trace the raw materials. Can trace main materials Can trace production date. Can't trace products Other Remark: according to purchase invoice. 			
4	Are corrective & preventive actions mechanism established and implemented effectively (including the suppliers/ sub-contractors control, incoming inspection, process control, final inspection and customer complaint)?	 Has the clear procedure and followed records Has the procedure but no written records. No written procedures or records. Other Remark: all actions were handled by general manager with suppliers on site. 			



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Part C: Product Research & Development Capacity

1.1 Product Research & Development Capacity	
The amount of R&D and relevant working experience.	 ☐ within 1year staff ☑ 6-10 years 1 staff ☑ 6-10 years 1 staff ☑ 0ver 10 years 1 staff Total 3 engineers
What is the main job responsibility for R&D engineers?	Design, Development of New Products, New technology; Declare patent certificate; Technology Support.
Is there any relevant design input, output, review, verification and validation documentation available for auditor to review?	No
Is there any special software or instrument used by the R&D staffs during the design process of new products? If yes, please list the main software or instrument.	
Does the company have an effective design change control procedure in place?	No Remark: all design change control procedure were controlled by R&D Manager
Please list the patent certificates and qualification license.	ZL 2012 1 0243549.4; ZL 20121 0029710.8



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Part E: Production Capacity & Quality Control

Remark: The audit content of this part relates only to its associated enterprise (Hubei Jiayu Better Organic Co., Ltd.)

Section 1: Production Capacity

1.1 Production Workflow Chart

				\$		
Raw Materials	12134 1761	Mixing & Mate	ching		Reaction	
				\$		
Cooling & Filling	ion (Top three	Function Test			Packaging Products	
Product	Price range	Min. order quantity	Top monthly output	Ave	erage monthly output	Total in 2016
API	Confidential	1 g	35 kg		9 kg	347 kg
Pharmaceutical Intermediate	Confidential	1 g	50000 kg	350	000 kg	420000 kg
1.3 Main Facilities			1			
Picture	Facility name		Brand or Country/Region of or	rigin	Target Value/machine*	Quantity
	Reaction Still		China		N/A	6 sets



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Measurement Tank	China	N/A	7 sets
Drying Machine	China	N/A	3 sets
Lab Test Still	China	N/A	16 sets
Rotary Evaporator	China	N/A	6 sets
Liquid Spectrometer Tester	China	N/A	1 set
Gas Spectrometer Tester	China	N/A	1 set
Ultrasonic Clean Tester	China	N/A	1 set



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Section 2: Production Process Control

2.1 Production Process Control		
ltem	Content	Observations /Comments
1	Product R&D capacity	 ☑ Own brand ☑ ODM ☑ OEM
2	Are the environmental conditions, such as tidiness and cleanliness being controlled and suitable for the operation performed?	 Very tidy Normal Need to improve Very poor
3	Are the necessary items /documents provided at appropriate location and under control?	 Work Instructions /procedures Workmanship standard /acceptance Golden sample /Approval sample Product picture Verbal by workshop director
4	Are written instructions available for incoming material inspections /testing? Is the relevant record maintained?	 Has instructions and uniformly followed Has instruction but no written records Materials checked by storage staff
5	Are written inspections /testing instructions available for finished products? Is the relevant record maintained?	 Have instructions and uniformly followed Have instruction but no written records Finished product checked by packing staff
6	What type of inspection is used for finished products?	 Random inspection Visual inspection Function inspection 100% inspection Visual inspection Function inspection
7	Are non-conforming units clearly marked/ segregated to prevent accidental dispatch?	 Marked and segregated Segregated but not marked clearly Not found on site
8	How are the non-conforming units handled?	 Repaired and re-inspection Picked out Used under control Others: Disused



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Part G: Working Environment

Section 1: Working Environment

1.1 Welfare Benefits			
ltem	Content	Observations /Comments	
1	Does the company have effective procedures to verify the age of staff at the time of recruitment?	 Has written procedure and keeps adequate age documents of workers Has written procedure but doesn't follow records Hasn't written procedure or follows records 	
2	Do all workers sign employment contracts with the factory?	 All workers sign employment contracts Some workers sign employment contracts Only management staff sign employment contracts No staff sign employment contracts 	
3	Is statutory contribution required for all employees' social insurance (e.g. health insurance, unemployment insurance, accident insurance etc.) paid for by the enterprise?	 All workers have social insurance. Some workers have social insurance Only management staffs have social insurance No staff have social insurance 	
4	Does the company have a clear and effective policy on working hours, rest and vacations? If does, please list it.	 All staffs work kept to the policy Most of the time it keeps to the policy except midseason. Usually needs overtime. No relevant records for working hours Describe the working hours: 	
5	Does the company pay extra remunerations for all overtime work?	 For all overtime work. For official holidays For official holidays except weekend. No extra remunerations for overtime. 	
6	Does the company have dormitories for staff? If yes, please describe the condition.	 Provide dormitories for all staff Provide dormitories for workers Provide dormitories for management staff No dormitories were provided. Describe the condition: 	
1.2 Labor	Protection		
ltem	Content	Observations /Comments	
1	Are there uniforms for all staff in company?	Yes No Other	
2	Is the emergency medical supplies enough and easily used in workshop?	☐ Yes ⊠ No ☐ Other	
3	Does the company arrange health and safety training for new workers?	Yes No Other	
4	Do the workers have the appropriate protective equipment during operation in workshop? Such as gloves, masks.	☐ Yes ☐ No ☐ Other	
5	Is there training needed and carried out for fire protection?	☐ Yes ☐ No ☐ Other	



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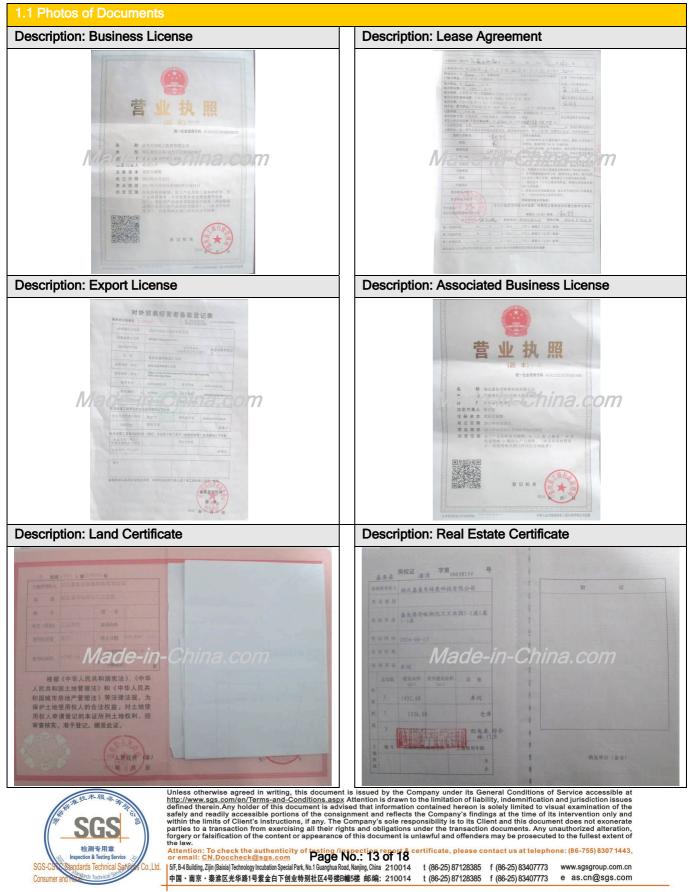
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Section 1: Photos of Documents



Part J: Photos





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Section 2: Photos of Company





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-- End of the Report --



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