



Supplier Assessment Report

Nanjing Aily Biotechnology Co., Ltd.

南京爱莉生物科技有限公司

This report is issued by Hangzhou Weiku Information Technology Co., Ltd. and Bureau Veritas Certification.

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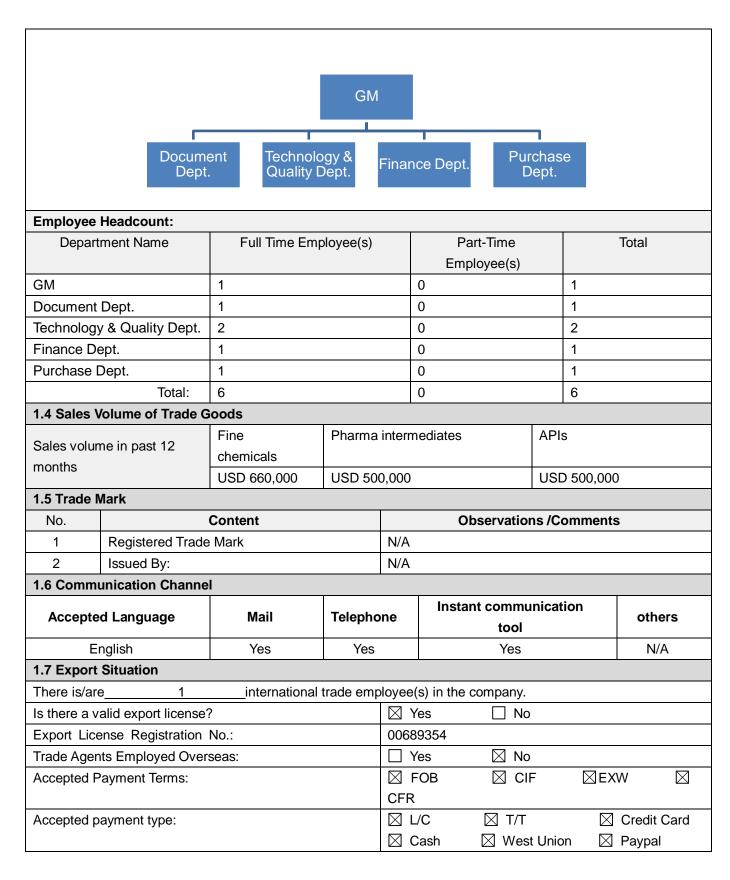
Section 1: Company Overview

Company Overview						
1.1 Legal Entity						
Does the supplier have the	□ Business license □					
following certificates to carry out	☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐					
business?	□ Organization code certificate					
Business license number:	320106000150973					
Validity period:	04/Sep./2009 03/Sep./2029					
Registered address:	Room 313, B Building, No. 26, Fuhougang, Gulou District, Nanjing City,					
registered address.	Jiangsu Province, China					
	Room 621, A2 Building, Science Park of Nanjing University of Science					
Company address:	and Technology, No. 129, Guanghua Road, Nanjing City, Jiangsu					
	Province, China					
Annual review conducted by:	Nanjing Industrial & Commercial Bureau Gulou Branch					
Name of legal representative:	Mr. Hongbin Chen					
Registered capital:	RMB 500,000. <u>00</u>					
Business type:	☐ Manufacturer ☐ Trading Company ☐ Both					
Type of ownership:						
	☐ Stated Owned ☐ Sole Proprietorship ☐ Other					
Products sold:	Fine chemicals, Pharma Intermediates, APIs					
Business contact person:	Mr. Hongbin Chen					
Phone number:	0086-25-83208295					
Fax number:	0086-25-83208202					
Email:	ben@ailychem.com					
Website address (URL):	http://ailybiotech.lookchem.com					
1.2 Company Facility Information						
Certification Type:	☐ Land Certification ☐ Real Estate Certificate					
	☐ Lease Agreement ☐ Factory Officer Claimed					
Number of Building(s)	1					
Total Size:	95 m ²					
1.3 Human Resources						
Company Chart						

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Lead time from order confirmation to production
delivery on average (products exiting the factory):

15 Days

1.8 Market Distribution (Previous 24 Months)

Products	Western Europe	Eastern Europe	North America	South Americ a	Asi a	Africa	Mid East	Domesti c
Fine	V	V	V	N	N	V	N	V
chemicals	٧	٧	٧	٧	V	٧	V	٧
Pharma	ما	ما		ما	a)	ما	2	2
intermediates	V	V	-	V	V	V	V	V
APIs	-	-	-	-	√	-	-	V

1.9 Certification & Photos

Business License (Duplicate) with Certificate / Records of Annual Inspection



Permit for Opening Bank Account

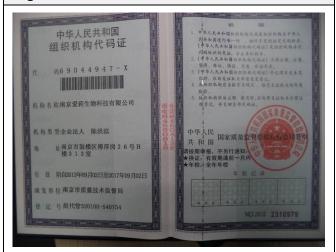


Custom Clearance Registration Form

Tax Registration Certificate



Organization Code Certificate



Import and Export Enterprise Registration







Hazardous Chemicals Business Permit





ISO9001:2008



1.10 Company Overview and Product Samples

Company Overview



Product Sample

Office Building



Product Sample







Product Sample





Product Sample

N/A





Section 2: Quality & After Sales and R&D Management

2.1 QA/QC and Purchasing Management									
	Content	Observations /Comments							
2.1.1	How many full time inspectors?	1							
2.1.2	Does the company have written inspection criterion?								
2.1.3	Does the company keep written inspection records?	☐ Comments: ☐ IQC ☐ IPQC ☐ FQC ☐ Comments:							
2.1.4	What is the type of finished products inspected?	 □ 100% of products with detailed inspection □ Random sampling inspection □ No inspection □ No inspection necessary 							
2.2 After Sa	lles Service								
	Content	Observations /Comments							
2.2.1	Are there any clear procedures for handling customer complaints?	 ☐ Yes, with clear procedures and written records ☐ Yes, with clear procedures but no written records ☐ Yes, with written records but no clear procedures ☐ No 							
2.2.3	Does the company have a written procedure for product alert and recall?	☐ Yes ☑ No							
2.3 R&D Ca	pacity								
	Content	Observations /Comments							
2.3.1	Number of special R&D engineer(s) in the company	1							
2.3.2	Does the company have a standard R&D procedure?	☐ Yes, with clear procedure and written instructions☐ Yes, with procedure but no written instructions☐ No							
2.3.3	Does the company have records of R&D processes?	☐ Yes ☐ No ☐ Comments: (typical process)							
2.3.4	Has the R&D product been internally tested and verified?	 ☐ Yes, with written record ☐ Yes, only partly have written record ☐ Yes, but no written record ☐ No 							

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	Were the R&D product inspected by the third	☐ Yes, all products were inspected by third party		
2.3.5	party organizations?	organizations		
2.3.3		☐ Yes, only partly		
		⊠ No		
	Is there a confirmation with client for the			
2.3.6	R&D product?	Yes, meet client requirements, but partly		
		confirmed		
		□ No		
2.3.7	If any, list all certifications and/or	N/A		
2.3.7	qualifications of the R&D department	N/A		
2.3.8	List special equipments for R&D purposes	Ultraviolet spectrophotometer, Moisture test		
2.3.0	List special equipments for R&D purposes	apparatus		

-- End of Report --

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