



Report Number: QIP-ASR2041181

**Audit Date** : 03 Sep.,2020

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 : M & G Products Co., Ltd.

镇江康如医疗器械有限公司

Showroom: <a href="http://mgproducts.en.made-in-china.com">http://mgproducts.en.made-in-china.com</a>

Address : Binjiang Avenue, LianheHuangshantao, Xinba Town, Yangzhong,

Zhenjiang, Jiangsu, China

Product : Syringe, Infusion Set, Surgical Tape, Bandage, Mask

#### has been on site audited for the Following Scope of Activity

1. General Information

- 2. Foreign Trade Capacity
- 3. Management System and Product Certification
- 4. Working Environment
- 5. Photos



#### **General Comments:**

M & G Products Co., Ltd. is a trading company of Syringe, Infusion Set, Surgical Tape, Bandage, Mask. It was established in 2013, located in Binjiang Avenue, LianheHuangshantao, Xinba Town, Yangzhong, Zhenjiang, Jiangsu, China. The company has passed ISO13485 medical device quality management system, and the products have passed CE certification. Products exported to Europe, South America other regions.

Sign for and on behalf of

SGS-CSTC Standards Technical Services Co., Ltd.



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SGS No: QIP-ASR2041181





# SUPPLIER ASSESSMENT REPORT

Audited Company	M & G Products Co., Ltd.
Audited Site:	Binjiang Avenue, LianheHuangshantao, Xinba Town, Yangzhong, Zhenjiang, Jiangsu, China

Consigner of Assessment	Made-in-China.com	4 I I I I I	
Audit Type	<ul><li>☑ Initial Audit</li><li>☐ Re-audit</li></ul>		
Audit Date	03 Sep.,2020	Verify Report	www.sgs.com/ecv
Auditor	Zhan Ning	Reviewed by	Toby Han

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# **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	91321182079852053Q
Year of established	25 Sep.,2013	Valid Date	Long Valid
Registered address	No. 968-970, Mingzhu Bay, Sanma	ao Street, Yangzhong,	Zhenjiang, Jiangsu, China
Actual address	Binjiang Avenue, LianheHuangsha Jiangsu, China	ntao, Xinba Town, Ya	ngzhong, Zhenjiang,
Does the company in abnormal operation status list of industrial and commercial bureau?	No		
Registered capital	RMB 800,000		
Name of legal representative	Mr. Yuan Ma		
Business scope	Sales of first class medical instrum and export business of all kinds of and acted as an agent (except the from import and export or manage subject to approval according to la approval by relevant departments)	of commodities and tech commodities and tech d by enterprises limite	echnologies is self-operated annologies that are prohibited d by the state). (For projects
1.2 Basic Information			
Contact person	Mr. Yuan Ma		
Phone number	0086-13705298675	Fax number 00	86-551-88324896
URL/Web address	http://mgproducts.en.made-in-china	a.com	
Company type		g Company Corporation	
Type of ownership	☐ Limited Company ☐ Public Company ☐ Foreign joint venture ☐ State-Owned ☐ Private Owner ☐ Wholly foreign-owned enterprises		
Associated company	Confidential		
Products manufactured / sold scope	Syringe, Infusion Set, Surgical Tap	e, Bandage, Mask	
1.3 Company Building Information			
According to:  Land certificate  Real estate certificate  Lease agreement  Observation Estimated on site			
The company area 300 square meter The land occupies 350 square meters	S.		
The offices occupy 300 square meter			
The workshops occupy 0 square meters.			



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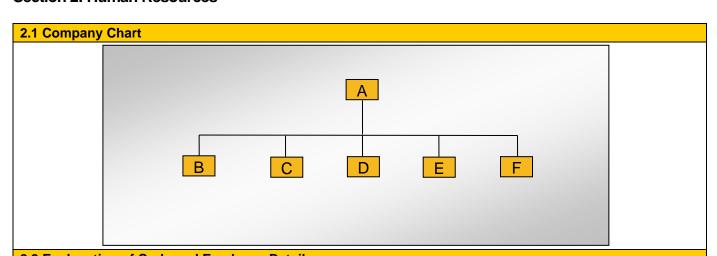
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#### **Section 2: Human Resources**



## 2.2 Explanation of Code and Employee Details

Code	Department	Number of employees
Α	General Manager	1
В	Financial Dept.	1
С	Purchasing Dept.	1
D	Sales Dept.	2
E	Admin Dept.	1
F	Quality Control Dept.	1
	Number in total:	7

# 2.3 Key Staff

Title	Full name	Education	Working experience for this trade / total experience
Confidential	Confidential	Confidential	Confidential



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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.	91321182079852053Q
The number of foreign trading staff with relevant trading experience.	☐ within 1 year staff ☐ 1-5 years 2 staffs ☐ 6-10 years staff ☐ over 10 years staff Total 2 staffs
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.	Confidential
Overseas agent / branch	☐ Yes
Nearest port	Shanghai 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)	<ul><li> within 15 workday</li><li> □ one month</li><li> □ 2-3 months</li><li> □ 4-6 months</li><li> □ 6-12 months</li><li> □ more time</li></ul>
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 Market	Main Product	Main client	
☐ North America			
South America	Syringe, Infusion Set, Surgical Tape, Bandage, Mask	Confidential	
□ Europe     □	Syringe, Infusion Set, Surgical Tape, Bandage, Mask	Confidential	
☐ Southeast Asia/ Mideast			
☐ Africa			
☐ East Asia (Japan/ South Korea)			
☐ Australia			
□ Domestic     □	Syringe, Infusion Set, Surgical Tape, Bandage, Mask	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?	<ul> <li>☐ Have the written procedures and followed records</li> <li>☐ Have the written procedures but no records</li> <li>☐ Have relevant records without written procedure</li> <li>☐ No written procedures or followed records</li> <li>☐ Other</li> </ul>		
2	Does the company have an updated list of approved suppliers?	<ul> <li>☐ The approved suppliers list was updated in 2020</li> <li>☐ Have the written suppliers without approved signature or date.</li> <li>☐ Provided some suppliers names</li> <li>☐ No approved suppliers list</li> <li>☐ Other</li> </ul>		

# **Section 4: After-sales Service Capacity**

4.1 After-s	.1 After-sales Service Capacity			
Item	Content	Observations /Comments		
1	Is there a procedure to conduct random product inspection after final packaging in place?	<ul> <li>☐ Have clear standards and written inspection records</li> <li>☐ No written standards but had inspection reports</li> <li>☐ Have the procedures but no inspection records</li> <li>☐ It's not necessary to carry out the inspection</li> <li>☐ Other</li> </ul>		
2	Is there a clear procedure for handling customer complaints?	<ul> <li>☐ Has the clear procedure and followed records</li> <li>☐ Has the procedure but no written records.</li> <li>☐ No written procedures or records.</li> <li>☐ Other</li> </ul>		
3	Can the finished/packaged product be traced by lot identification to the appropriate raw materials test reports?	<ul> <li>☑ Have the procedures to trace the raw materials.</li> <li>☑ Can trace main materials</li> <li>☑ Can trace production date.</li> <li>☐ Can't trace products</li> <li>☐ Other</li> </ul>		
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)?	<ul> <li>☐ Has the clear procedure and followed records</li> <li>☐ Has the procedure but no written records.</li> <li>☐ No written procedures or records.</li> <li>☐ Other</li> </ul>		



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# **Part D: Management System and Product Certification**

1.1 Management System and Product Certification	
Management system certification	☐ Others     Certificate Name: ISO13485:2016     NO.: SX601283830001     Issued by: TUV Rheinland     Issued Date: 29 Jun.,2018     Valid Until: 12 Jun.,2021     Scope: Manufacture and Distribution of Medical Devices
Product certification	☐ CE Certificate Name: CE NO.: DD60148986 Product Name: Medical Devices Model: N/A Standard: 93/42/EEC Issued by: TUV Rheinland Issue Date: 08 May.,2020 ☐ UL
	□ ccc
	□ RoHS
	☐ Others
Test reports for raw materials	RoHS
	Reach
	☐ Others
	⊠ NIL



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# **Part G: Working Environment**

## **Section 1: Working Environment**

1.1 Welfa	re Benefits	
Item	Content	Observations /Comments
1	Does the company have effective procedures to verify the age of staff at the time of recruitment?	<ul> <li>☐ Has written procedure and keeps adequate age documents of workers</li> <li>☐ Has written procedure but doesn't follow records</li> <li>☐ Hasn't written procedure or follows records</li> </ul>
2	Do all workers sign employment contracts with the factory?	<ul> <li>✓ All workers sign employment contracts</li> <li>☐ Some workers sign employment contracts</li> <li>☐ Only management staff sign employment contracts</li> <li>☐ No staff sign employment contracts</li> </ul>
3	Is statutory contribution required for all employees' social insurance (e.g. health insurance, unemployment insurance, accident insurance etc.) paid for by the enterprise?	<ul> <li>✓ All workers have social insurance.</li> <li>☐ Some workers have social insurance</li> <li>☐ Only management staffs have social insurance</li> <li>☐ No staff have social insurance</li> </ul>
4	Does the company have a clear and effective policy on working hours, rest and vacations? If does, please list it.	<ul> <li>✓ All staffs work kept to the policy</li> <li>✓ Most of the time it keeps to the policy except midseason.</li> <li>✓ Usually needs overtime.</li> <li>✓ No relevant records for working hours</li> <li>Describe the working hours: 8 hours per day, from Mon. to Fri.</li> </ul>
5	Does the company pay extra remunerations for all overtime work?	<ul> <li>☐ For all overtime work.</li> <li>☐ For official holidays</li> <li>☐ For official holidays except weekend.</li> <li>☐ No extra remunerations for overtime.</li> </ul>
6	Does the company have dormitories for staff? If yes, please describe the condition.	<ul> <li>□ Provide dormitories for all staff</li> <li>□ Provide dormitories for workers</li> <li>□ Provide dormitories for management staff</li> <li>☑ No dormitories were provided.</li> <li>Describe the condition: N/A</li> </ul>
1.2 Labor	Protection	
Item	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 The company is a trader and does not engage in production activities
3	Does the company arrange health and safety training for new workers?	☐ Yes☐ No☐ Other The company is a trader and does not engage in production activities
4	Do the workers have the appropriate protective equipment during operation in workshop? Such as gloves, masks.	☐ Yes ☐ No ☑ Other The company is a trader and does not engage in production activities



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production activities	5	Is there training needed and carried out for fire protection?	<ul> <li>☐ Yes</li> <li>☐ No</li> <li>☑ Other The company is a trader and does not engage in production activities</li> </ul>
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### Part J: Photos

#### **Section 1: Photos of Documents**

#### **Description: Business License**



#### **Description: ISO13485**



#### **Description: Export License**



#### **Description: EC**





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### **Section 2: Photos of Company**

**Description: Company Gate** 



**Description: Product Samples** 



**Description: Product Samples** 



**Description: Office** 



**Description: Product Samples** 



**Description: Product Samples** 





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