



Report Number: QIP-ASI205150

Audit Date : 19 Mar.,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 : Qingdao Miuton Medical Co., Ltd.

青岛明药堂医疗股份有限公司

Showroom : http://qdmiuton.en.made-in-china.com

Address : 2F/3F/4F, A1, No.1, Jinye Road, High-Tech Industrial Development

Zone, Qingdao, Shandong, China

Product : Face Mask, Alcohol Wipes, Wet Wipes, Medical Sterilizing Ultrasound

Couplant, Fever Cooling Patch, Band-aid, Sterilized Cotton Ball, PVPI

Cotton Ball, Liquid Wound Dressing, Medical Nursing Pad

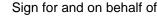
has been on site audited for the Following Scope of Activity

1. General Information

- 2. Foreign Trade Capacity
- 3. Product Research & Development Capacity
- 4. Production Capacity & Quality Control
- 5. Working Environment
- 6. Energy Saving and Emission Reduction
- 7. Photos

General Comments:

Qingdao Miuton Medical Co., Ltd. is a manufacturer with 100 employees; it was established in 2004, located in 2F/3F/4F, A1, No.1, Jinye Road, High-Tech Industrial Development Zone, Qingdao, Shandong, China; The offices and workshops occupy an area of 3394 square meters; Qingdao Miuton Medical Co., Ltd. has rich experience in domestic trade.



SGS-CSTC Standards Technical Services Co., Ltd.



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SUPPLIER ASSESSMENT REPORT

Audited Company	Qingdao Miuton Medical Co., Ltd.
Audited Site:	2F/3F/4F, A1, No.1, Jinye Road, High-Tech Industrial Development Zone, Qingdao, Shandong, China

Consigner of Assessment	Made-in-China.com		
Audit Type			
	☐ Re-audit		
Audit Date	19 Mar.,2020	Verify Report	www.sgs.com/ecv
Auditor	Wayne Wang	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?		Registration Number	91370200760288916E	
Year of established	17 May.,2004	Valid Date	Long Valid	
Registered address	2F/3F/4F, A1, No.1, Jinye Road, H Shandong, China	ligh-Tech Industrial D	evelopment Zone, Qingdao,	
Actual address	2F/3F/4F, A1, No.1, Jinye Road, H Shandong, China	ligh-Tech Industrial D	evelopment Zone, Qingdao,	
Does the company in abnormal operation status list of industrial and commercial bureau?	No			
Registered capital	RMB 37,318,000			
Name of legal representative	Mr. Yuqiang Li			
Business scope	Production, sales: Class I medical materials, sanitary supplies, import need to be authorized by the law c department).	and export of goods a	nd technology. (Items which	
1.2 Basic Information				
Contact person	Ms. Qianqian Zhi			
Phone number	0086-13626480815	Fax number Nil		
URL/Web address	http://en.miuton.com			
Company type		Company Corporation		
Type of ownership	☐ Limited Company☐ Public C☐ State-Owned☐ Private		n joint venture foreign-owned enterprises	
Associated company	No			
Products manufactured / sold scope Face Mask, Alcohol Wipes, Wet Wipes, Medical Sterilizing Ultrasound Couplant Fever Cooling Patch, Band-aid, Sterilized Cotton Ball, PVPI Cotton Ball, Liquic Wound Dressing, Medical Nursing Pad			lizing Ultrasound Couplant, , PVPI Cotton Ball, Liquid	
1.3 Company Building Information				
According to: Land certificate Real estate certificate Lease agreement Observation Estimated on site				
The company area 3394 square meters The land occupies square meters. Confidential The offices occupy 594 square meters. The workshops occupy 2800 square meters.				



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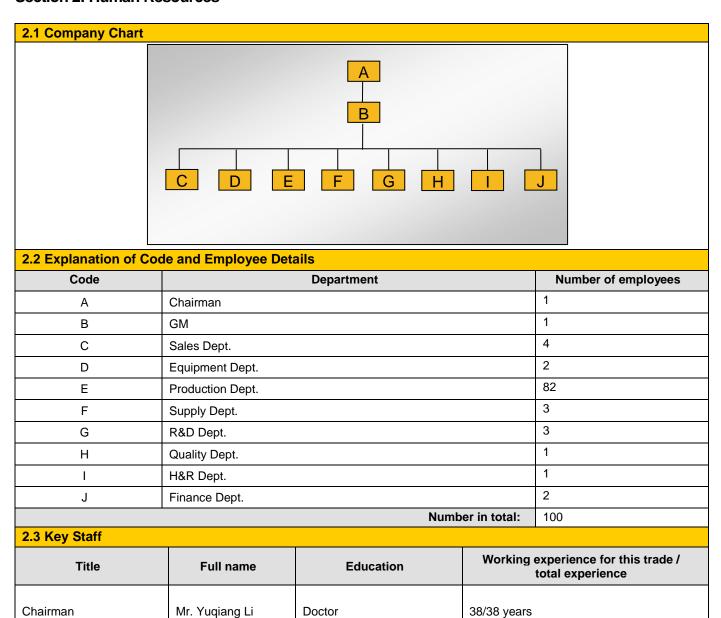
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Section 2: Human Resources



Remark: Above information of Key Staff was based on interview with 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.	91370200760288916E
The number of foreign trading staff with relevant trading experience.	☐ within 1 year staff ☐ 1-5 years staff ☐ 6-10 years staff ☐ over 10 years 3 staffs Total 3 staffs
The language freely used by foreign trade staff	
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	Qingdao Port
Acceptable quotation terms	☑ FOB ☑ CIF ☑ CFR
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				
☐ South America				
☐ Europe				
☐ Southeast Asia/ Mideast				
☐ Africa				
☐ East Asia (Japan/ South Korea)				
☐ Australia				
□ Domestic □	Face Mask, Alcohol Wipes, Wet Wipes, Medical Sterilizing Ultrasound Couplant, Fever Cooling Patch, Band-aid, Sterilized Cotton Ball, PVPI Cotton Ball, Liquid Wound Dressing, Medical Nursing Pad	Confidential		
Others				



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Section 3: Supplier Management

3.1 Supplier Management			
Item	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	
2	Does the company have an updated list of approved suppliers?	 ☑ The approved suppliers list was updated in 2019 ☐ Have the written suppliers without approved signature or date. ☐ Provided some suppliers names ☐ No approved suppliers list ☐ Other 	

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?	 ☐ 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 		
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 		
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 		
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 		



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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 ☐ 1-5 years staff ☐ 6-10 years staff ☐ over 10 years 3 staff ☐ Total 3 engineers
What is the main job responsibility for R&D engineers?	Researching and developing of new products and providing technical support.
Is there any relevant design input, output, review, verification and validation documentation available for auditor to review?	YES
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.	YES instrument: HPLC, High Efficiency Gas Chromatograph
Does the company have an effective design change control procedure in place?	YES
Please list the patent certificates and qualification license.	patent certificates: ZL 2012 2 0132585.9, ZL 2012 2 0183433.1, ZL 2012 2 0183339.6, ZL 2014 2 0068984.2, ZL 2011 2 0228716.9, ZL 2012 1 0091868.8, ZL 2012 1 0004607.8, ZL 2012 1 0020699.9, ZL 2015 1 0066045.3, ZL 2015 2 0694237.4, ZL 2015 2 0089677.7, ZL 2016 3 0478880.3; qualification certificates: Medical Device Production License: 20120071



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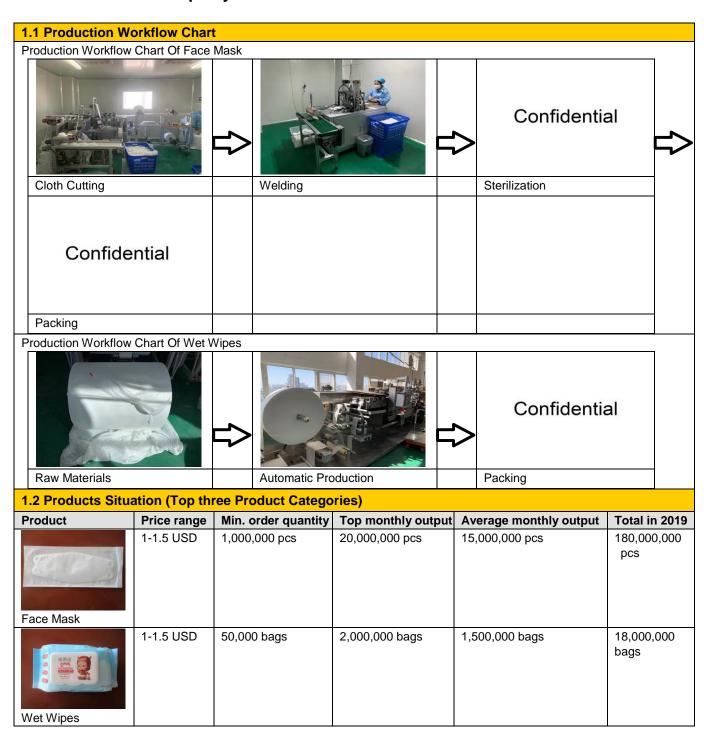






Part E: Production Capacity & Quality Control

Section 1: Production Capacity





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	1-1.5 USD	100,000 boxes	300,000 box	es	150,000 boxes	1,800,000 boxes
Fever Cooling Patch						
1.3 Main Facilities						
Picture	Facility name	Country/Regio	on of origin		Value/machine*day	Quantity
	Wet Wipes Production Lir	China ne		Confide	ential	3
	Face Mask Production Lir	China		Confide	ential	3
	Alcohol Wipes Production Machine	S China		Confide	ential	1
	Water Genera	ttor China		Confide	ential	1
	Blending Mac	hine China		Confide	ential	2



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Section 2: Production Process Control

2.1 Pro	2.1 Production Process Control			
Item	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 		



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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?	 ☐ 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: 8 hours per workday
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: Nil
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
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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Part H: Energy Saving and Emission Reduction

Section 1: Environmental Management

1.1 Environmental Management			
Item	Content	Observations /Comments	
1	Environmental Management System	☐ ISO 14001 Certificate☐ Cleaner production management☑ NIL	
2	Environmental Impact Assessment	☐ The Report of Environmental impact assessment is approved by EPA .☑ NIL	
3	Check and Acceptance of Completed Constructive Projects	 ☐ The Report Completed Constructive Projects Inspection and Acceptance is approved. ☑ NIL 	
4	"Three Simultaneity" for Environmental protection	☐ The Report of "Three Simultaneity" for Environmental protection Inspection and Acceptance is available.☒ NIL	

Section 2: Water, Gas and Noise Control

2.1 Water, Gas and Noise Control		
Item	Content	Observations /Comments
1	Water-intaking	☐ Water-intaking license☐ No water-intaking license☒ No water taken from natural water body
2	Waste water	 □ Water-draining license or contract □ Having carried out waste water treatment before draining □ Having carried out waste water monitoring and the result of which being within the standard limit ☑ No detail activities were found
3	Emission to air	 ☐ Air pollutant emission license ☐ Having carried out air pollutant emission monitoring and the result of which being within the standard limit ☒ No detail activities were found to reduce emission to air
4	Classification and Recycling for solid waste	 ☐ Having applied trash classification recycling ☐ Having special warehouse for hazardous waste ☐ No detail activities were found
5	Noise	 ☐ The company carried out noise monitoring and the result of which being within the standard limit one time per year ☒ No detail activities were found



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Part J: Photos

Section 1: Photos of Documents



Description: Medical Device Production License



Description: Certificate of Invention Patent



Description: Export License



Description: Trademark



Description: Certificate of Utility Model Patent





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Section 2: Photos of Company

Description: Company Building



Description: Workshop Inside



Description: Lab



Description: Office



Description: Lab



Description: Lab





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Description: Product Sample



Description: Product Sample



Description: Product Sample



Description: Product Sample



Description: Product Samples



Description: Product Samples



-- End of the Report --



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