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Explanation about the Vendor Assessment Methodology

The Vendor Assessment was based on the three broad parameters i.e., Physical Location (Company Profile), Financial Capacity and Production Capacity.

The data regarding company profile and financial capacity of the vendor were auto-fetched from the GeM Portal, through their registration process. QCI validated the quality certificate, test reports and business experience of the vendor through the documents submitted by the official on behalf of the firm. An undertaking has been taken by the vendor to verify the debarred/blacklisted status of the firm. ITR verification was done by GeM through the PAN details provided by the vendor. The production capacity was verified based on the documents received by the firm.

The documents are considered to be authentic and complete, based on vendor's claims and cannot be verified for any misrepresentation, concealment or forgery. The documents such as process flow chart, type of machines, safety measures, transportation etc. provided by the firm were validated through a video-based assessment and geo-tagged pictures.

It is clarified that the assessment was based on mutual trust between parties. QCI stands guarantee to the data collected and verified exclusively at the time of assessment. However, if at a later date, it is found that any vendor has purposefully submitted documents which were either incorrect, modified, misrepresented in order to gain unlawful benefits, the GeM will be free to take any action as deemed under law.

In case, the video assessment was inconclusive, an on-site assessment will be conducted to factually check the authenticity of the capability, as claimed. The vendor may be asked to pay for the expenses of this verification assessment.





1. Authentication of Vendor Profile

Parameters	ers Requirements Responses		
	Name of the Company/Firm	EVALYN HEALTHCARE SOLUTIONS PRIVATE LIMITED	
	Registered Address	Office No A 503, Plot No 59, Navi Mumbai, Mumbai, Thane, Thane, MAHARASHTRA, 400614	
	Address of Manufacturing Site	1F, 3F & 4F, No. 23, Wu Quan Road, New Taipei, Industrial Park, Wu Ku District, New Taipei City - 24886, Taiwan	
	Primary Contact Number	9967674411	
	Primary Email ID	harishjagdale@evalynhealthcare.com	
Company Description/Details	Additional Contact Number	9967674411	
	Additional Email ID	info@evalynhealthcare.com	
	Constitution	company	
	Company Identification Number	U24100MH2022PTC382869	
	Date of Incorporation of the Company	18/05/2022	
	GST Number (Company)	27AAHCE0466E1ZT	
	PAN of the Company	AAHCE0466E	
	Company debarred/blacklisted by any Govt. organisation	No	
	Bank Name	ICICI BANK LIMITED, KHARGHAR, NAVI MUMBAI	
Banking Details	Account Holder Name	EVALYN HEALTHCARE SOLUTIONS PRIVATE LIMITED	
	Account Number	103005002905	
	IFSC code	ICIC0001030	
Details of Products	Quality Certification	Yes	
Registered for VA	Other License Number (if applicable)	No	





Details of top three Government & Private Orders (Value wise) received in last three financial years:

A) Government Organisation

S.No.	S.No. Ordering Govt. Item Name Department/Agency		Quantity Supplied	Order Value (Rs.)	Work Order/Completion Letter Number
-	-	-	-	-	-

B) **<u>Private Organisation</u>**

S.No.	Ordering Entity	Item Name	Quantity Supplied	Order Value (Rs.)	Work Order/Completion Letter Number
-	-	-	-	-	-

2. Financial Capacity of the Vendor

Financial Capacity: Annual Turnover of the Firm with Profit/Loss Statement of Last Three Years.

S.No.	Acknowledgement No.	Assessment Year	Turnover (in Rs.)	Profit/Loss Amount (in Rs.)	ITR Type
-	-	-	-	-	-





3. Production Capacity of the Vendor

Parameters	Requirements	Responses	
Basic Information	Number of production sites	1	
	Generic 'Process Flow Chart' availability	Yes	
Process Capability	Availability of 'Specific Process Flow Diagram' of the products manufactured by the firm	Νο	
Production	List of all the machines/software involved in the manufacturing process	Plastic Molding Machine, Sealing Machine, etc.	
Capacity	Firm's monthly manufacturing capacity	500 Nos	
	Utilisation of the capacity against the manufacturing capacity	0 Nos	
	Customisation according to customer specifications	No	
Quality	Availability of 'Quality Assurance Plan' document	No	
Quanty	Availability of 'Part-Wise Inspection Plan'	Νο	
Territory of	Territory of Operations	N/A	
Operations	Availability of SOP for defect free delivery of products	No	
Safety	Availability of 'Safety Standard and Operating Procedures' manual	Νο	
	List of the approved suppliers along with quantity supplied	Yes	
Suppliers	Availability of document for performance monitoring of suppliers	Νο	
Rejected Material	Process of handling rejected material	The rejected material is disposed.	
Consumer	Availability of consumer complaint process	Νο	
Grievance	Time taken for resolution of complaints (in days)	-	
	Availability of warranty/ guarantee or recalling to the product	Νο	
Warranty	Obligation for raw material, product development design, product quality, product lifetime and product sustainability in the warranty provided by the firm	Νο	
	Availability of in-house company R&D centre	No	
	Availability of R&D centre of the parent company	Νο	
	Recognised or approved R&D centre	Νο	
Research and Development	Collaboration with premium R&D institutes for product development/new product/quality improvement	Νο	
	Patents/Copy Rights/ published/quality improvement	Νο	
	Prototypes Developed	No	





Items for which vendor assessment is requested

5.No	Category Name	Production Capacity (per month)	Test Reports available with the firm	Status
1	poct point of care test and bio markers	500 Nos	No	OEM
	sessment Feedback			
	as a Quality Certificate. for the product is not available.			
e vendor d	pes not have any 'Past Work Experience'.			





4. Video Assessment

Parameters	Information	Responses
Physical Location	Physical location verified with the address proof provided in the Desktop Assessment	Yes
Basic Information	Name of the official taking part in the assessment	Mr. Clas Sivertsen
	Designation of the official taking part in the assessment	Founder and Chief Science Officer
Process Capability	Process flow for production followed by the firm	Completely Followed
Production Capacity	Machines available for production	All machines available
Quality	Quality Assurance Plan' being followed by the firm	Yes
Suppliers	Key aspects for selecting suppliers	Quality and time.
Territory of Operation	Availability of transport facility	Yes
Safety	Presence of Safety Signs	Yes
	Availability of 'Safety Equipment'	Yes
Research And Development	Availability of R&D facility	No
	Products/Processes available for which patents have been obtained	No
	Availability of developed prototypes	No

Video Assessment Feedback

1) Based on the interview with Mr. Clas Sivertsen (Founder and Chief Science Officer) along with the live video, pictorial evidence, and geo-tag verification, Evalyn Healthcare Solutions Private Limited is found to be Deemed OEM of Amedifact Co. Ltd.

2) Necessary raw materials such as LED camera, circuit board, mother board were evidenced at the assessed site.

3) The machines such as electric screwdriver, pneumatic Insertion machine & PCB carrier were evidenced during the video assessment.

4) The quality check process involves functional test of the finished product.











Physical Location: Outside of the Firm

Raw Material Stock



Production Process



Critical Machine for Production







Quality Check Process



Safe Packaging for Defect Free Delivery of Products



Safety Equipment



Finished Product

Recommendations on the basis of Vendor Assessment

The firm is a Deemed OEM





IMPORTANT NOTICE

Quality Council of India's Vendor Assessment Report is a confidential, third party assessment, on an entity desirous of claims of OEM status under GeM and is solely intended for the specific recipient to whom it is addressed. It cannot be disclosed to any other person, authority or organisation without specific consent of GeM or the entity, which is assessed.

The Vendor Assessment Report is based on the information/clarification provided to QCI by the management/officials of the concerned entity through self-disclosure, which is presumed to be honest, unambiguous and correct. QCI doesn't guarantee or take responsibility for the accuracy, completeness or adequacy of any information on which report is based unless verified by QCI in Vendor Assessment. QCI accepts no liability for the contents of this report or for the consequences of any actions taken on the basis of information provided in the report. Vendor Assessment is not an audit and this report is not in any way a recommendation for entering into any kind of transaction/commitment/agreement etc. with the concerned entity.

QCI accepts no responsibility, whatsoever, for any loss or damage from the use of any information given in the report and nothing contained in the report is capable or intended to create any legally binding obligations, financial or any other liability whatsoever on QCI, it's leadership or persons/organisations involved in Vendor Assessment.

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