ACPL

Quotation Request Form

Important: For preparation of a written quotation, we need information about your organization. All information supplied by you will be treated in strict confidence. Please complete this questionnaire. Use extra sheets wherever required. Fields marked with "* "are mandatory for filling.

			COMP	ANY DE	TAILS				
*Company Name:									
* Registered Address:									
*Site Address: (Temporary)									
Phone:			F	ax:					
*E-mail:	Website:								
*Chief Executive/MD:			N	/lobile:					
*Contact Person Name:			Р	osition:		N	/lobile:		
Company Status (Please Tick): Limited Liability Partnershi		olic Lim Other P	nited Priv	ate Lim	nited □	Partners	hip □ Prop	rietary	
Please list the number of employees in each area/site (please use additional sheets if required)	Full Time	Part Time	Contract Employees	Shifts	Full Time (Site 2)	Part Time (Site 2)	Contract Employees (Site 2)	Shifts (Site 2)	Personnel working away from the premises
Manufacturing/Service area									
Quality Control/Technical									
Administration									
Storage/Warehouse									
Other									
Management									
Total Employees (Full time equivalent)									
Total no of employees doing repetitive jobs Employees directly involved in scope of management system QMS:, EMS:, OH&SMS:, FSMS:, ISMS:, MD-QMS:, EnMS:, ABMS Note: If more than one site, please give address/details on back of this page. No of Temporary Sites (In operation at present)									
		(CERTIFICAT	ION/S F	REQUEST	ΓED			
Certification Required (Please ISO 22000:2018 ☐ ISO 27000:2018 ☐ ISO 27000:2018 ☐ ISO 27000:2018 ☐ ISO 27000:2019 ☐ ISO 2700	27001:2 n ロ R	2022 le- Cer	□ ISO 134 tification □	85:201 Transfe	6 🗖 er Certifica	ISO 500 ation from	01:2018 [other CAB		7001:2016
Quality Management Systems Number of Sites to be Audited Is there any process that affect Other Exclusions, If any Legal Obligations if any Whether company is responsite Is the Clause" Design & Development Systems Legal Obligations of the Clause of t	d? □ Si cts the p ble for lopmen	ngle Coroduct demon t" inclu	Multiple conformity a stration of proded in the So	oduct/so	ervice per	— formance		No	
□ Environmental Managemental Number of Sites to be Audited Whether Initial Environmental Whether Register of Significant Whether Legal Register availant Whether Environmental Management Has EMP been implemented?	d? □ Ši Reviev nt Aspe able? □ agemen	ngle L v (IER) cts / Im I Yes L t Progr	I Multiple available? pacts availal I No am (EMP) av	☐ Yesble?		l No s □ No			

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CDI		01-1	D	-
ACPL		Quotation	Request	Form
□ Occupational Health & Sa Number of Sites to be Audited Have you identified Hazards? Detail all identified Critical occ Whether Incident/ Accident Re Imp: Please furnish form 03 Q	? ☐ Single ☐ Multiple ☐ Yes ☐ No upational health and safe egister available? ☐ Yes I RF Annexure- OH&SMS	ety risks □ No	-	
Attached as above □ Yes □ N □ Food Safety Management		3		
Number of Sites to be Audited Have you implemented HACC Any seasonality issues? Total No of HACCP Studies (August 1984) How many process lines are the seasonal to the seasona	? P Principles? As per ISO/TS 22003:201	13)	☐ Single ☐ Mu☐ Yes ☐ No☐ Yes ☐ No☐ Yes ☐ No☐ O	ultiple
Any Prior Audits Conducted If Yes, attach audit findings	·		□ Yes □ No	
Other Factors(Kindly Confire Product Types=; Produ Building Area=; Infraste □ Information Security Mana	uct Lines= ; Productructure= ; In House	e Lab Testing=		
Has a Statement of Applicabili No. of user = No. of servers = Any Prior Audits Conducted f Yes, attach audit findings:		No. of sites = No. of Workstatic ∕es □ No	ns (PC + Laptop	
Factors related to business and	organization (other than			
Category Type(s) of business and regulatory requirements	☐ Organization works in nor ☐ Organization has custome ☐ Organization works in cri	ers in critical business se	s and non-regulated se	ectors ^a
	Standard processes with			
Process and tasks	workunder the organization's Standard but non-repetitiv Complex processes, high r of certification (ISMS covers	ve processes, with high nu number of products and s	same tasks; few produ Imber of products or se ervices, many busine	cts or services ervices ss units included in the scope
Process and tasks Level of establishment of the MS	work under the organization's Standard but non-repetitiv Complex processes, high r	control carrying out the reprocesses, with high nu number of products and s highly complex processe ished and/or other mana nanagement systems are	same tasks; few products or so ervices, many busine is or relatively high nugement systems are in implemented, others	cts or services ervices ss units included in the scope imber or unique activities) n place not

Category	Grade			
ITT's Country of the second of	☐ Few or highly standardized IT platforms, servers, operating systems, databases, networks, etc.			
IT infrastructure complexity	☐ Several different IT platforms, servers, operating systems, databases, networks			
	☐ Many different IT platforms, servers, operating systems, databases, networks			
Daniel den en e	☐ Little or no dependency on outsourcing or suppliers			
Dependency on outsourcing and suppliers, including cloud services	\square Some dependency on outsourcing or suppliers, related to some but not all important business activities			
	☐ High dependency on outsourcing or suppliers, large impact on important business activities			
Information System development	☐ None or a very limited in-house system/application development			
	☐ Some in-house or outsourced system/application development for some important business purposes			
	☐ Extensivein-houseoroutsourcedsystem/application development for important business purposes			

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Medical Device Quality Management System ISC			
Number of Sites to be Audited? Outsourced process:	Single □ Multiple		
Critical activity:			
Question		Yes	No
Is the product a nearly finished and assembled medic	cal device? (i.e., it is intended to be used for	100	110
a medical purpose and only needs packaging and/or			
Is the product intended to be a component/part of a n			
Is the organization contracted to carry out any activiti			
regulation (e.g., relabeling, remanufacturing of other	medical devices)?		
Is the product supplied sterile?			
Does the product contain software developed by the			
Is "Design and Development" in the scope of the ISO permits exclusion of design and development which i			
devices)?	s the case very often for low-risk medical		
Is the product (Raw Materials, Parts, Components, S	ubassemblies Maintenance Services or		
Other Services) intended to support associated media			
Note: Refer to the note in Annex A, Table A.1.7, a) as			
*Kindly select applicable answer in above question se	eries.		
□ Energy Management System ISO 50001:2018			
Number of Sites to be Audited? ☐ Single ☐ Multiple			
Annual Energy Consumption=			
Number of energy Sources= Number of significant energy uses (SEUs) =			
	16		
□ Anti-Bribery Management System ISO 37001:201 Number of Sites to be Audited? □ 3	<u>ro</u> Single □ Multiple		
	Yes No		
	Yes □ No		
In Case of Integrated Management Systems, Kindly		vel of Inte	egration in
Integration	%		. J
1. An integrated documentation set, including WIs to a good level	I of development, as appropriate; □		
Yes □ No 2. Management Reviews that consider the overall business strate	gy and plan D. Van. D. No.		
	gy and plan do res do No		
 3. An integrated approach to internal audits □ Yes □ No 4. An integrated approach to policy and objectives □ Yes □ N 	0		
5. An integrated approach to systems processes □ Yes □ No			
6. An integrated approach to improvement mechanisms, (C			
measurement and continual improvement); and, Yes			
7. Integrated management support and responsibilities. Yes	□ No		
Other Certification Program Requested (Number of Sites to be Audited? Single Multiple) Any Prior Audita Conducted	l Voc El í	No
Number of Sites to be Addited? • Single • Multiple	Any Prior Audits Conducted □ If Yes , att		
Accreditation: ☐ ACCREDITATED ☐ NON ACC		aon addi	timanige
Scope for Certification:	REDITATED		
Scope for Certification:			
RUS	INESS DETAILS		
Identify products / services of your company	INCOO DE TAILO		
racinally products a convictor of your company			
Activities being performed outside the main site:			
(i.e. activities at temporary sites e.g. construction, collection of samples, service	ce delivery etc.)		
Outsourcing if any:			
Outsourcing it arry .			
Name of the Consulting Organization:			
Identify key processes in manufacturing or provision of		ontrol,	
Purchasing, Marketing/Sales, Maintenance, Stores, HR	D etc)		
Any statutory & regulatory requirements related to Prod			
GST No: TIN No_			
PAN No CIN No			
Main Customers: Main Supp	liers:.		

Declaration: The information provided above is true to the best of our knowledge and behalf.

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ACPL	Quo	tation Request Form	
Quotation Requested by	Name:		
Designation:	Sign:	Date:	
Reviewed By :	FOR THE USE OF AC	Date:	
Can this Application be further prod	cessed □ Yes □ No		

Please send it on below address or Email:

ACME CERTIFICATION PVT. LTD.

2-A/3, SECOND FLOOR (FRONT PORTION), ASAF ALI ROAD, TURKMAN GATE, NEW DELHI-110002 Ph: +91 9811010507, Email: ewsdelhi@rediffmail.com, Web: www. Acmeregistrar.com

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