QEHS		Request for Proposal cum Application Form							
1. Organization De	tails								
Name of org. *:									
Register Address 1 *:									
Working / HO address: (For Multi-site, fill annexure 01)									
Contact Person*:					Dept./	Position:			
Mob. /Tel.*:					Fax:				
Website:					E-Mai	 * •			
	ard *.				L-IVIGI				
Require ISO Standa	aru ·:	T			1				
☐ ISO 9001	ISO14	001	☐ OH&S 450	001 S2000] ISO 27001	ISO 20000-1	Otl	ners
Mode of Audit	Mode of Audit O		Virtual/Online-audit Combine Audit (Onsite & Online Audit				ine Audit		
	<u> </u>		The type of plat	form to be ι	used i.	e., Google m	eet/ Zoom:		
Certification Program Required * Initial Surveillance Recertification Transfer Scope Expansion									
1.1 Existing Certific	cation (If	fapplic	able)						
ISO 9001						ners			
Name of Certification Body: Certification No: Expiry Date:									
If Other, please spe	ecify:								
Scope applied *:									
(i.e., Manufacture	of								
abc)									
2. Number of Emp	loyees *								
Activities		Shifts	Full Time F		Perfor type o	ming Same of Job	Temporar Unskilled workers	y <u>I</u>	Effective No. of Employees, Filled by QEHS
Management									
Production Area									
Quality Control/Technical									
Administration		\bot							
Other									
Total No. of Employees									
Operation for Weekend / weekly holiday:									
3. Organization Activity									
Legal Status of Organization (i.e., Proprietor/ Pvt. Ltd. Or Partnership) *:									
Applicable Legal, statuary & regulatory act:									
(i.e., Labour law)									
Organization Key Process Area:									

(i.e., purchase/store/production etc.)						
Organization Products/Services:						
(i.e., abc & xyz products etc.)						
Any outsourcing process:						
(i.e., printing etc.)						
Is the organization part of some larger						
organization: if Yes (Please furnish the name)	∐ Yes					
0.8a2a.a.a.	∐ No					
Language of document and record:	Language	of audit:				
4. Operation condition of Management System*						
1. Has the documented management system bee	n operated	and maintained?	Yes No			
- Approval/Implementation Date of Policy:						
- Approval/Implementation Date of Manual, Pro	cedure and	Instruction:				
2. Is the Internal Audit carried and the effectivene	ss of the au	dit is confirmed?	Yes No			
- Date of Internal Audit:						
3. Is management review carried?			Yes / No			
- Date of management review:			,			
4. If operating multi-site, you			N/A Yes (Filled			
are same corporate?			Annexure 01)			
are same corporate:are actually implementing same activity in cor	trol2		Yes No			
(3) have same CEO?	iti Oi :					
0 (,					
(4) are using same quality system and procedure		/	☐ Yes ☐ No			
* Please attach detail information about multi-site	(Total site	no./size and location of each	☐ Yes ☐ No			
site)						
4.1 This section Applicable for		ntal Management System ISC	J 14001:2015			
How many Sites the company is Managing at the s						
Do you have Register of Significant Environment aspect? Yes No						
Do you have An Environmental Management Man Do you have An Internal Environmental Audit Prog		☐ Yes ☐ No				
4.2 This section Applicable for Occupa						
Hazard's Identified? Yes No						
nazaru s identinied: Tes INO						
Detail any critical occupational health & safety risks identified?						
4.3 This section Applicable fo	r Food Safe	ty Management System ISO	22000:2018			
Number of Sites to be Audited?		Single Multipl	e			
Have you implemented HACCP Principles?		Yes No				
Any seasonality issues?		☐ Yes ☐ No				
If yes, give detail on separate sheet.	242)					
Total No of HACCP Studies (As per ISO/TS 22003:20 How many process lines are there in production						
FSSAI License Registration No:	r					
Any Prior Audits Conducted		☐ Yes ☐ No				
If yes, attach audit findings						
Other Factors (Kindly Confirm No's): -						
Product Types=; Product Lines=; Product Development=; CCP=; OPRP=; Building						
Area=; Infrastructure=; In House Lab	Testing=	; Translator Requiremer	nts= ;			
4.4 This section Applicable for Inf			ISO 27001:2013			
1. SOA Version No	D	ate of implement:				

	s (i.e., IT / Data Centre/Server) sessment & Risk Treatment, If yes on which date
4. Business complexity [Ple	ease select type of complexity in your org. as per A, B, C, chose any no. from 1 or 2 or 3]
A. Type(s) of business and regulatory requirements	 ☐ 1. Organization works in non-critical business sectors and non-regulated sectors; ☐ 2. Organization has customers in critical business sectors; ☐ 3. Organization works in critical business sectors.
B. Process and tasks	 ☐ 1. Standard processes with standard and repetitive tasks; lots of persons doing work under the organization's control carrying out the same tasks; few products or services ☐ 2. Standard but non-repetitive processes, with high number of products or services ☐ 3. Complex processes, high number of products and services, many businesses units included in the scope of certification (ISMS covers highly complex processes or relatively high number or unique activities).
C. Level of establishment of the MS	 ☐ 1. ISMS is already well established and/or other management systems are in place ☐ 2. Some elements of other management systems are implemented, others not ☐ 3. No other management system implemented at all, the ISMS is new and not established.
5. IT complexity [Please sel	ect type of complexity in your org. as per A, B, C, chose any no. from 1 or 2 or 3]
A. IT infrastructure complexity	 ☐ 1. Few or highly standardized IT platforms, servers, operating systems, databases, networks, etc. ☐ 2. Several different IT platforms, servers, operating systems, databases, networks ☐ 3. Many different IT platforms, servers, operating systems, databases, networks.
B. Dependency on outsourcing and suppliers, including cloud services:	 ☐ 1. Little or no dependency on outsourcing or suppliers ☐ 2. Some dependency on outsourcing or suppliers, related to some but not all important business activities ☐ 3. High dependency on outsourcing or suppliers, large impact on important business activities.
C. Information System development:	 □ 1. None or a very limited in-house system/application development □ 2. Some in-house or outsourced system/application development for some important business purposes □ 3. Extensive in-house or outsourced system/application development for important business purposes.
6. Confirmation of access to	o organizational records
audit team. 2. Not agreed to s review by audit team,	all the ISMS records or information about design and effectiveness of controls for review by hare all the ISMS records or information about design and effectiveness of controls for because it contains confidential or sensitive information. 3", please mention the information which cannot be shared in during the audit:
	e for Information Technology - Service Management System requirements ISO 20000-1:2018 ighly standardized IT platforms, 1 or 2 servers, operating system, database, networks
2. Several different IT	platforms, 3-5 servers, operating system, database, networks Database,
Is there any ITSMS Docume	ntation & records which cannot be made available for review by the audit team because they sitive information and to provide the corresponding justification.

Kindly provide list of such information:
4.6 This Section is applicable for Medical Device Quality Management System ISO 13485:2016
1. Number of Sites to be Audited?
2. Any Outsourced process:
3. Critical activity:
4.7 This section is applicable for Learning Services Management System
Number of Sites to be Audited? Single Multiple
Methodology of Learning is Described Yes No
Any conflicts regarding Learning Services Management system
Outsourced process:
5. Details of the consultancy firm/consultant has taken consultancy for ISO 9001/ISO 14001/ISO 45001/ISO 22000/ISO
27001/ISO 20000-1 or other standard process in organization:
1. Consulting Company Name:
2. Write the name of consultant:assisted for establishment of system.
6. Other details of Organization
Note: if required, please attach relevant supporting document(s)
DECLARATION*: The above information is true to the best of my knowledge and belief and I am authorized to provide such
information on behalf of the above said organization, we request an estimate/Quotation as above mention. In case of
multi-site, kindly fill annexure 01.
Person Name (who Filled Application):
This section is for Office (QEHS) Use only.
Acceptance, Ref. No Not Acceptance
Supplement information is needed:

Note: "*" fields are mandatory.

Annexure -01 (For Multi-site)

Site Country Name:						
Name of Site*:						
Address 1 *:						
Contact Person*:				Dept./Position:		
Mob. /Tel.*:				Fax:		
Website:			E-Mail*:			
Require ISO Standard *:				1		
Site Scope/Activity:						
(i.e., Manufacture of						
abc)						
2. Number of Employees:						
Activities	Shifts	Full Time	Part time	Performing Same type of Job	Temporary Unskilled workers	Effective No. of Employees, Filled by QEHS
Management						
Production Area						
Quality Control/Technical						
Administration						
Other						
		Total No	o. of Employee	es		
Operation for Weekend,	/ week	ly holiday:				

In case, if you have more site's, kindly fill this annexure per site.