

Application Submission & Review:

• Submit your application to help us understand your requirements.

Stage 1 Audit (Document Review):

 Our auditors evaluate your organization's documentation, policies, and processes against the requirements of the standard.

Stage 2 Certification Audit:

 Our auditors verify the implementation and effectiveness of your management system against the requirements of the standard

Corrective Actions (if necessary):

 If any non-conformities are identified during the audits, you'll have the opportunity to address them

Certificate Release:

• IAS grants Certification post successful audit.

ABOUT THE CGMP STANDARD

cGMP is a globally recognized set of regulations and guidelines for ensuring the quality, safety, and consistency of pharmaceutical, biotechnology, and medical device products. It provides a systematic framework for manufacturing processes that meet stringent quality standards. cGMP focuses on key principles:



QUALITY MANAGEMENT SYSTEM



DOCUMENTATION AND RECORDS



PERSONNEL AND TRAINING



FACILITIES AND EQUIPMENT



PROCESS CONTROLS AND VALIDATION



COMPLAINT HANDLING AND PRODUCT RECALL

Elevate your manufacturing quality with cGMP, ensuring products are consistently produced and controlled according to quality standards, while minimizing risks to patients and consumers.



ACHIEVE EXCELLENCE WITH INTEGRATED ASSESSMENT SERVICES



WHY CHOOSE IAS?



Our skilled auditors have over 15 years of industry experience to provide valuable insights during the certification process.



We tailor our services to meet your specific needs, ensuring a seamless certification journey.



We are with you beyond certification, assisting with maintenance and updates as your business evolves.



Our certification services are internationally recognized, giving you a competitive edge.





Unlock Success with Integrated Assessment Services:
Your Path to Certified Excellence!



• Opportunities for business growth

and market credibility through

high-quality products