imarc
WE'LL EARN YOUR APPROVAL.

AUDITING

With regulatory inspections on the rise, you need to be sure your site adheres to the most stringent requirements for compliance and patient protection. Our rigorous auditing process is designed to **identify and address issues before the FDA finds them and prepare your team to answer questions with confidence.**

Our comprehensive auditing services are based on the FDA's BIMO checklists and cover every aspect of your study, including protocol, IRB requirements, agreements, and regulations. In addition to GCP audits, we also provide vendor qualification audits and clinical quality audits against standard operating procedures.

There are unique risks to consider at every stage of a clinical trial. That's why we use a multi-level approach to prepare your team for approval that includes:

1. EVALUATING YOUR SITES AND VENDORS
2. INSPECTING YOUR FACILITIES, DATA AND PROCESSES
3. PREPARING YOUR STAFF FOR INSPECTIONS AND INTERVIEWS

Benefits of IMARC Auditing:

- Assess performance against research requirements
- Ensure the quality and integrity of data collected
- Identify negative process-level trends
- Receive meaningful deliverables about your study, including a summary of findings, an assessment of your greatest strengths, weaknesses and opportunities, and expert recommendations to achieve compliance