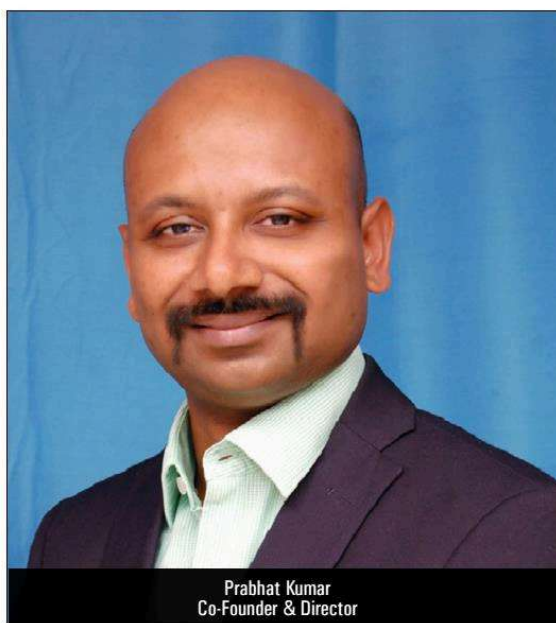


CLINTASK

TRANSFORMING COMPLIANCE AND QUALITY MANAGEMENT WITH TECHNOLOGY-DRIVEN EFFICIENCY AND BESPOKE SOLUTIONS



Dr Mahesh Kumar
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The clinical trial services market is evolving rapidly, driven by technological advancements, personalized medicine, and globalization. India has emerged as a key player, particularly in clinical data management and safety case processing, where its combination of technology and human resources excels.

However, challenges remain. India's clinical trial site infrastructure lags behind the U.S., Europe, China, and South Korea. Over the past decade, the industry has transitioned from paper-based processes to technology-driven systems. This shift, further accelerated by the pandemic, underscores the growing demand for speed and efficiency in trials, especially for database setups and regulatory approvals.

India's diverse patient pool and cost-effective solutions make it an attractive destination for global trials. The rise of digital transformation and hybrid trial models positions the country as more than just an outsourcing hub, it's becoming a global leader in clinical trial services.

To this end, Clintask is at the forefront of quality management, QA, computerized system validation (CSV), and compliance services for life sciences, pharmaceuticals, and biotech industries. The company differentiates itself by offering highly customized solutions, focusing on GxP auditing (GLP, GCP, GMP, GVP) and emerging compliance needs like GDPR and data privacy.

Unlike traditional providers, Clintask emphasizes agility and a purpose-driven approach, avoiding

bureaucratic delays common in larger organizations. This client-centric strategy allows them to deliver tailored solutions, meeting diverse needs across sectors from small biotech startups to large pharmaceutical companies and IT service providers.

Moreover, the team is uniquely positioned, working across diverse sectors, including small biotech, large pharmaceutical companies, CROs, and IT service providers. This broad experience equips Clintask to offer solutions that are not only compliant with global standards like FDA and EU regulations but also adaptable to the specific needs of each client. Its ability to avoid bureaucratic delays, typical of larger organizations, ensures a more agile, purpose-driven approach. This flexibility, combined with a commitment to understanding each client's unique needs, has earned the company a growing client base, with over 15 clients in the US alone.

"By focusing on purpose-driven solutions rather than rigid templates, we stand out as a highly adaptable and client-focused partner in the clinical trials industry", adds Dr Mahesh Kumar, Co-Founder & Director, Clintask.



WHAT SETS CLINTASK APART IS ITS ABILITY TO DELIVER PRAGMATIC AND FLEXIBLE SOLUTIONS TO CLIENTS OF ALL SIZES, FROM LARGE MULTINATIONAL PHARMA COMPANIES TO EMERGING BIOPHARMACEUTICALS AND E-CLINICAL TECHNOLOGY FIRMS

Founders' Vision and Expertise

Founded by Prabhat Kumar and Dr Mahesh Kumar, Clintask was built on a foundation of transparency and trust. With over 150 regulatory inspections under their belt, the founders bring a wealth of experience across global markets, including USFDA, EMA, MHRA, and PMDA.

Their entrepreneurial journey started with outreach to trusted colleagues and mentors, resulting in long-term engagements. Today, Clintask counts top global pharmaceutical companies among its clients, including a leading U.S. based firm in the top 15 globally and a top five India based pharma company.

This focus on personalized, pragmatic solutions over generic templates has driven the company's growth, solidifying its reputation as a reliable partner.



Ambitious Roadmap for the Future

Clintask has set its sights on becoming one of the top five global providers of Quality and CSV services within the next three years. Despite being a team of 20, the company leverages technology to enhance service efficiency, aiming to reduce client quality assurance costs by up to 30 percent.

Currently, 80 percent of Clintask's clients are based in the U.S., with the remainder in Europe and Asia. The company is expanding its global footprint, focusing on agile auditing, data privacy assessments, and regulatory inspection support.

In early 2024, the company showcased its commitment to sustainability by presenting a case study on sustainable quality assurance at the Society of Quality Assurance (SQA) in the U.S. This initiative aligns its services with global sustainability goals, modernizing traditional quality management approaches.

Client Success and Impact

Clintask's client-centric approach is evident in its collaborations. For a small biotech company, the company quickly implemented a risk-based QMS tailored to their needs, streamlining compliance and ensuring smooth trial operations.

The company's expertise in CSV and data privacy assessments ensured that the client's systems, including ePRO/eCOA and EDC, were fully compliant and operational from the outset, minimizing delays and safeguarding patient data.

This combination of technical expertise and personalized service has made Clintask a trusted partner for startups and multinationals alike, driving strategic growth in the pharmaceutical sector. "With our comprehensive vision, we are keen to lead in transforming quality management into a strategic, sustainable, and scalable service", concludes Prabhat Kumar, Co-Founder & Director, Clintask. **P0**