COVID-19 Clinical Trial Analysis Report

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Executive Summary

This report analyzes the **current status and key findings** of COVID-19 clinical trials, focusing on **recruitment trends, regulatory approvals, and market readiness**. Key observations include:

- Half of number of trials were in the recruiting phase, screening participants for eligibility.
- Remdesivir and convalescent plasma therapy received marketing approval in April 2023.
- 100 pediatric trials were actively enrolling participants across 50+ locations.
- Trial durations vary significantly, with most spanning **2 to 12 months**, but some extending **beyond 100 months** for long-term safety monitoring.
- 50% of trials were progressing through Phase 2 & Phase 3, critical steps toward regulatory approval.
- **22 Phase 4 trials** completed post-market surveillance in **2020–21**, assessing long-term efficacy.
- 56 Phase 3 trials wrapped up between 2020–21, paving the way for marketing approval.
- Behavioral trials on Facebook Ads saw the highest enrollment, with the "Importance of Staying Safe on Thanksgiving".

Introduction

This study aims to analyze the **status and findings** of COVID-19 clinical trials, providing insights into **recruitment trends**, approved treatments, trial durations, and regulatory readiness.

Methodology

- Data was sourced from Covid-19 Trails Data Excel file.
- Checked & Analyzed for Missing Values, Removed Duplicates and set desired datatypes wherever required using **Python(Pandas)**.
- Exported the Excel file as **Covid-19 Trails(cleaned)**.
- **Tableau** was Used for Data Visualization and fetching Key insights.

Key Findings

Recruitment Trends

- 48.50% of trials are in **recruitment**, actively screening patients.
- Early-phase trials are enrolling **faster**, while late-stage trials face **delays may be due to regulatory approvals.**

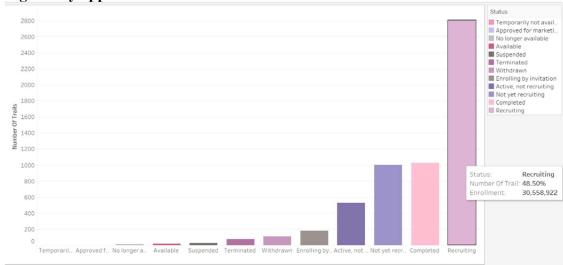


Figure 1 - Shows the Percentage of Trail in Recruiting Status

Approved Treatments

- **NCT04338360** Convalescent Plasma Therapy is using plasma from recovered patients to boost immune response got marketing approval in April 2023.
- NCT04323761 Remdesivir (Antiviral Drug), an antiviral drug showing effectiveness in severe cases received approval for marketing in April 2023.

Pediatric Clinical Trials

- Around 82% of trials set an age criterion above 17 years, limiting pediatric research.
- 100 active pediatric trials focusing on ages 1 month 17 years on Over 50 locations conducting interventional studies for children.

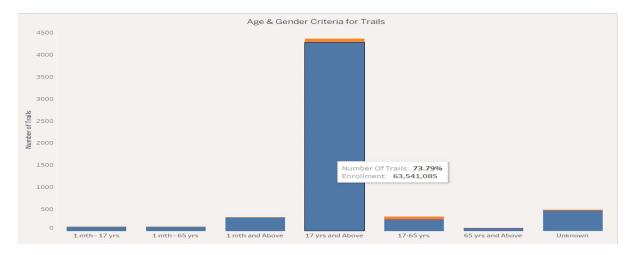


Figure 2 - Maximum Percentage of Age Group for Number of Trails

Trial Durations

- Majority span 2 to 12 months, but some exceed 100 months for extended monitoring majority of which is Observational Trails.
- Extended trials track COVID-19's mutational behavior, immunity sustainability, and long-term health implications.

Phase 2 & 3 Trials

- 50% of trials were currently in Phase 2 & 3, key stages for efficacy assessment.
- Phase 2 trials involve small patient groups, focusing on dosage optimization and initial effectiveness checks.
- Phase 3 trials expand to larger populations, gathering comprehensive safety and efficacy data before regulatory submission.

Post-Market Surveillance

- 22 Phase 4 trials completed between 2020–21, now undergoing long-term safety monitoring.
- Phase 4 studies assess the real-world impact of approved treatments, refining recommendations for dosage and usage.
- These studies track rare side effects, interactions with other medications, and overall effectiveness beyond initial trials.

Regulatory Approvals & Market Readiness

- **56 Phase 3 trials** wrapped up between **2020–21**, paving the way for **marketing approval** after successful regulatory evaluation.
- Regulatory bodies assess **patient outcomes**, **safety profiles**, **and trial efficacy** before approving treatments for large-scale distribution.
- Approved treatments are expected to be integrated into global COVID-19 management strategies.

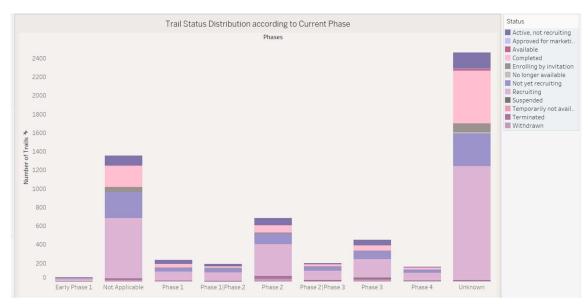


Figure 3 - Number of Trails according to Current Phase and Status

Behavioral Trials & Digital Outreach

- The highest enrollment in behavioral trials was recorded in the Facebook Ads campaign—"Importance of Staying Safe on Thanksgiving," with 2 million participants joining.
- This highlights the power of social media-driven health awareness in engaging mass audiences as Digital contribution is not just time saving but easy to access compared to traditional outcomes.
- Digital outreach for behavioral studies proves effective in **public health messaging**, vaccine acceptance, and preventive strategies.
- This detailed breakdown **ensures clarity, depth, and impact** while maintaining **logical structure**.

Conclusion

The findings highlight **both progress and challenges** in COVID-19 treatment development, emphasizing **recruitment trends**, **regulatory approvals**, **and post-market surveillance**. The high engagement in **behavioral trials via digital platforms** suggests **new opportunities** for public health outreach, demonstrating the growing influence of **technology-driven health campaigns**.

As we move forward, the **Digital Revolution** holds **immense potential** for enhancing interventional trial surveillance, enabling real-time tracking, improved patient engagement, and more efficient data collection. Leveraging **AI** and digital tools could further streamline clinical research, ensuring faster, more accessible medical advancements in the near future.

Thank you for reviewing this report. I appreciate your time and consideration.

Best Regards,

Samiran Bhagat