DSRB GREEN LANE

Accelerating Clinical Trial Reviews Through Collaborative Excellence

Version 1.1, Dated 28 Jul 2026

Our Holistic Green Lane Approach

DSRB **Green Lane** provides comprehensive end-to-end IRB support from start to finish.

- **Priority queue** processing for faster timelines
- **Pre-submission reviews** to optimise your application
- **IRB quality assurance**
- **Post-approval monitoring** for ongoing study success.

We partner with you through **fast-track** ethical reviews to arrive at **prompt outcomes** for your clinical trials.

DSRB Green Lane FAQs

Is the Green Lane for everyone?

- No, Green Lane is crafted to support industry sponsored clinical trials only.

Does Green Lane guarantee approval?

- No. Studies still undergo the same ethical and regulatory review standards. It supports submission readiness and smoother review processes but does not influence the review outcomes.

How does it help accelerate timelines?

- The service is intended to streamline submission preparation and reduce avoidable revision cycles by identifying submission issues early, improving application completeness.

What support does the Green Lane come with?

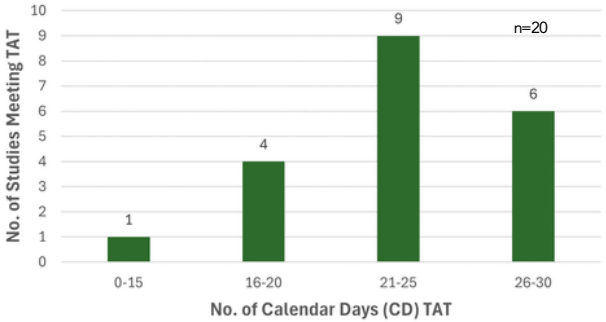
- Escalation to Board Reviewers for pre-Full Board reviews.
- Administrative guidance on submission requirements, documentation completeness, workflow coordination, and communication support throughout the review process.

Can Green Lane support amendments or renewals?

- Yes, the Green Lane service provides IRB support throughout the lifecycle of the study including amendments and renewals.

1 PRIORITY IRB REVIEW
Fast, predictable review for eligible Green Lane studies

PRIORITY QUEUE FOR NEW STUDIES



23 DAYS
Average Time Taken from Submission to Outcome*

100%
Reviewed in ≤ 30 Calendar Days

100%
Single IRB Tabling

*Data based on outcome date Nov24 – Jun26

Special Acknowledgement to NHG Health Domain C Chairperson and Board Members

3 POST-APPROVAL MONITORING
Supporting your study throughout the lifecycle for ongoing study success

OUR PERFORMANCE

Amendments Review	Study Renewals	Safety Events Review^
16 DAYS Average Turnaround Time*	27 DAYS Average Turnaround Time*	09 DAYS Average Turnaround Time*
53 Reviewed	10 Reviewed	104 Reviewed
89% Reviewed Within 30 Calendar Days	100% Renewed before expiry	100% Reviewed Within 30 Calendar Days

*Data based on outcome date Dec24 – Jun26

^Safety Events includes Deviation/Non-Compliance (DNC), Severe Adverse Events (SAE), Unanticipated Problem Involving Risks to Subjects or Others (UPIRTSO) Reports

2 COLLABORATIVE PRE-SUBMISSION SUPPORT
Helping you submit with confidence

HOW WE SUPPORT YOU

Dedicated IRB Team
A responsive partner throughout your study.

Protocol and ICF Screening
We provide guidance on IRB requirements and offer relevant insights.

Full Board Preparation
We engage our IRB members early to prepare for Full Board (FB) review.

Query Resolution
FB queries are posted within 5 work days. Analysts will actively engage the study team to resolve queries promptly.

4 QUALITY ASSURANCE
Continuous quality improvement for trustworthy reviews

COMPREHENSIVE AUDIT FRAMEWORK

Green Lane reviews undergo independent quality checks.

What Sponsors Say

★★★★★
Analysts are excellent, approachable and professional. Responses are prompt and consistent.

★★★★★
Always responsive, provides clear feedback, and handles tight timelines with ease. I appreciate this premier service a lot!

★★★★★
I would like to acknowledge the assistant analyst for his dependable support with admin matters. His prompt clarifications have been a big help.

SPONSOR CONFIDENCE
100%
Based on post-review surveys

Call to action

Partner us. Position your clinical trial on the DSRB Green Lane.

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Your Trial Matters To Us