





RIDERSTM

The Research & Innovation Document Engagement and Review Service

Enhance Your Clinical Trial Documents with

Expert Review

- Streamline Ethical Review Process and Improve Participant Understanding
- Expert reviews of Participation
 Information Sheet (PIS) and informed
 Consent Forms (ICF).
- A service aligned with **HRA guidance**.
- Competitive pricing in line with NIHR National Costing and Validation Review (NCVR).





Recommendations Report

"The ability to discuss our requirements with expert advisors in RIDERS prior to submission has been incredibly helpful in ensuring that we have everything prepared correctly"

- Feedback from one of our commercial partners







- **Comprehensive review** covering ethical, legal, and study management aspects.
- Fast turnaround to meet deadlines.
- Access to experienced/ diverse and experienced Patient and Public Involvement (PPI) groups.
 - Valuable feedback to improve participant understanding.

