Welcome to the Clinical Data Research Hub (CDRH), a coordinating centre for clinical research, hosted by the Liggins Institute and supporting the ON TRACK Network. In collaboration with the Department of Obstetrics and Gynecology, and Department of Paediatrics: Child and Youth Health, University of Auckland.

We provide a comprehensive range of data management services for randomised trials and clinical studies, including online databases, custom randomisation, drug distribution and web dashboards for study monitoring. Our systems will help you streamline setting up and running your trial.

Contact: researchhub@auckland.ac.nz  +649237713
Randomisation and databases: www.ligginstrials.org

**CDRH Guides**
- Planning a Trial
- Clinical Trial Terminology
- Study Protocol
- Ethics
- Trial Committees
- Authorship
- Data Sharing
- Mori Responsiveness
- Standard Operating Procedures

**Randomisation**
- CDRH Randomisation Site
- CDRH Study Numbers
- Randomisation Plans

**Data Management**
- REDCap FMHS
- REDCap Liggins
- REDCap Resources
- Variable Naming Convention
- Study Reports

**Analysis**
- Normal Distribution Calculator
- OpenEpi - Open Source Statistics
- Medcalc Calculators
- SAS Guides
- NZDep2013 Index of Deprivation
- Trial Analysis
- Data Cleaning
- Diagnostic Tests

**Follow-up**
- Bayley Scales Age Calculator
- LifePath Follow-up SharePoint
- NZ Travel Calculator
- Definitions for Follow-up Studies
- Middlemore Parking and Clinic Bookings
- WHO Child Growth Standards (0-5 years)
- WHO Growth Reference 2007 (5-19 years)

**Consensus Statements and Checklists**
- EMA Guidance for COVID-19 Pandemic
- Equator Network
- INQUire Framework for Evaluating Research Quality

**Protocols**
- SPIRIT 2013 Statement
- SPIRIT 2013 Explanation and Elaboration
- ACTA 2019 Implementability

**Outcomes**
- COMET - Core Outcome Measures in Effectiveness Trials
- COS - Prevention of Preterm Birth
- Neonatal Infection Case Definition (Brighton Collaboration)

**Reporting**
- CONSORT 2010 - Reporting Parallel Group RCTs
- CONSORT 2010 - Extension to Cluster RCTs
- CONSORT 2010 - Extension to Noninferiority / Equivalence RCTs
- SIRONG Checklist - Reporting of Neonatal Nutrition and Growth
- STROBE Statement - Reporting Observational Studies
- STROBE Statement - Explanation & Elaboration
- STARD 2015 - Diagnostic Accuracy Studies
- TIDieR Checklist - Reporting of Interventions
- Very Preterm Birth Studies (Pediatrics 2016)

**Data & Safety Monitoring**
- Guidelines for the Content of Statistical Analysis Plans (JAMA 2017)

**Laboratory**
- Liggins Institute Analytical Capabilities

**Training**
- NIH Protecting Human Subjects certificate
- NIH GCP course
- ROCHE GCP course (runs best in Firefox)
- Cultural Competency
- Infection control
- Definition of Research and Audit
- Classification of Observational Studies
- Standford Medical Statistics Certificate

**Key Links**

**ANZ**
- Australia & New Zealand Clinical Trials Register (ANZCTR)
- Health and Disability Ethics Committees NZ (HDEC)
- Health Research Council NZ (HRC)
- IMPACT Clinical Trials Network
- MedSafe NZ
- Neonatal Unit Contacts NZ
- Online Forms NZ (HDEC / SCOTT)
- DHB Research Contacts
- Therapeutic Goods Agency (TGA) Australia
- Kidz First Neonatal Research
- Cure Kids Portal
- Ministry of Health NZ
- ACC Act 2001: Clinical Trials
- NEAC Ethical Standards 2019

**International**
- Cochrane Neonatal
- Cochrane Pregnancy
- Cochrane Childhealth
- Global Obstetrics Network (GoNET)
- ICH GCP
- NIH Clinical Research Toolkit
- PROSPERO
- Trial Forge UK
CDRH Admin

- Redmine
- CDRH SharePoint (runs best in IE)
- Liggins VPN Access
- University of Auckland Staff Intranet
- Space statistics
- LiFePATH Admin
- CDRH Randomisation Demo (code 991000tst)
- CDRH Data Environment

Synthesis & Translation

- PRISMA-P 2015 - Preferred Reporting for Systematic Review and Meta-analysis Protocols
- PRISMA 2009 - Preferred Reporting Items for Systematic Reviews and Meta-analyses
- PRISMA-IPD 2015 - Preferred Reporting Items for a Systematic Review and Meta-analysis of Individual Participant Data
- PRISMA Extension 2015 - Network Meta-analyses
- PRISMA-Č Extension 2016 - Systematic Reviews and Meta-analyses of Newborn and Child Health Research
- PRISMA-DTA - Preferred Reporting Items for a Systematic Review and Meta-analysis of Diagnostic Test Accuracy Studies
- MOOSE - Reporting of Meta-analysis of Observational Studies in Epidemiology
- GRADE - Grading Quality of Evidence and Strength of Recommendations
- GRADE - Non-Randomised Studies
- GRADE Handbook
- Cochrane Handbook 2019 - Systematic Reviews of Interventions
- AGREE II - Assessing Quality and Reporting of Practice Guidelines
- PROSPERO Registration of Systematic Reviews - Checklist
- RoB2 - Cochrane Risk of Bias Tool for RCTs
- ROBINS-I - Risk of Bias In Non-randomized Studies of Interventions
- Risk of Bias Tool for Observational Studies
- Meta-analysis of Animal Studies
- Median to Mean Calculator
- WHO Guideline to Rapid Reviews

• DAMOCLES - Charter for Clinical Trial Data Monitoring Committees
• EROIN 2011 - Standards Requirements for GCP-Compliant Data Management in Multinational Clinical Trials
• NAESS 2019 - Neonatal Adverse Event Severity Scale (INC)

• Trial Managers Network
• NLM Journal Database