

Multicenter Clinical Trials: Leveraging Network Infrastructure to Advance Research and Outcomes for Women, Children, and Pregnant and Lactating Individuals

NACHHD June Council 2022

Bob Tamburro, MD, MS on Behalf of the Division of Extramural Research







Amniocentesis: HEW Backs Test for Prenatal Diagnosis of Disease

The study, which was conducted by researchers at nine major medical centers,* was coordinated and supported by the National Institute of Child Health and Human Development (NICHD). The study was designed to answer two basic questions about the use of amniocentesis during the middle 3 months of pregnancy.

7 NOVEMBER 1975

Initiative Background

- NICHD has long supported multicenter clinical studies
- NICHD formed its first clinical research networks in 1985
 - Neonatal Intensive Care Units
 - Maternal-Fetal Medicine Units
- Perform large scale clinical trials
- Network support resulted in clinical practice changes



^{*}The participating institutions were: Children's Memorial Hospital, Chicago; Eunice Kennedy Shriver Center, Boston; Johns Hopkins University School of Medicine, Baltimore; Mount Sinai School of Medicine, New York; University of California at Los Angeles-Harbor General Hospital, Torrence; University of California School of Medicine, San Diego; University of Michigan School of Medicine, Ann Arbor; University of Pennsylvania School of Medicine, Philadelphia; Yale University School of Medicine, New Haven.

The New England Journal of Medicine

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Volume 329 OCTOBER 21, 1993 Num

PREVENTION OF PREECLAMPSIA WITH LOW-DOSE ASPIRIN IN HEALTHY NULLIPA

Reduction of Maternal-Infant Transmission of Human Immunodeficiency Virus Type 1 with Zidovudine Treatment

Edward M. Connor, Rhoda S. Sperling, Richard Gelber, Pavel Kiselev, Gwendolyn Scott, Mary Jo O'Sullivan, Russell VanDyke, Mohammed Bey, William Shearer, Robert L. Jacobson, Eleanor Jimenez, Edward O'Neill, et al., for the Pediatric AIDS Clinical Trials Group Protocol 076 Study Group*

Low-dose aspirin for the prevention of preterm delivery in nulliparous women with a singleton pregnancy (ASPIRIN): a randomised, double-blind, placebo-controlled trial

ORIGINAL ARTICLE

A Trial of Hyperimmune Globulin to Prevent Congenital Cytomegalovirus Infection

B.L. Hughes, R.G. Clifton, D.J. Rouse, G.R. Saade, M.J. Dinsmoor, U.M. Reddy, R. Pass, D. Allard, G. Mallett, L.M. Fette, C. Gyamfi-Bannerman, M.W. Varner, W.H. Goodnight, A.T.N. Tita, M.M. Costantine, G.K. Swamy, R.S. Gibbs, E.K. Chien, S.P. Chauhan, Y.Y. El-Sayed, B.M. Casey, S. Parry, H.N. Simhan, P.G. Napolitano,

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JAMA | Original Investigation

Effect of Therapeutic Hypothermia Initiated After 6 Hours of Age on Death or Disability Among Newborns With Hypoxic-Ischemic Encephalopathy

A Randomized Clinical Trial

Abbot R. Laptook, MD; Seetha Shankaran, MD; Jon E. Tyson, MD, MPH; Breda Munoz, PhD; Edward F. Bell, MD; Ronald N. Goldberg, MD; Nehal A. Parikh, DO, MS; Namasivayam Ambalavanan, MD; Claudia Pedroza, PhD; Athina Pappas, MD; Abhik Das, PhD; Aasma S. Chaudhary, BS, RRT; Richard A. Ehrenkranz, MD; Angelita M. Hensman, MS, RNC-NIC; Krisa P. Van Meurs, MD; Lina F. Chalak, MD, MSCS; Amir M. Khan, MD; Shannon E. G. Hamrick, MD; Gregory M. Sokol, MD; Michele C. Walsh, MD, MS; Brenda B. Poindexter, MD, MS; Roger G. Faix, MD; Kristi L. Watterberg, MD; Ivan D. Frantz III, MD; Ronnie Guillet, MD, PhD; Uday Devaskar, MD; William E. Truog, MD; Valerie Y. Chock, MD, MS-Epi; Myra H. Wyckoff, MD; Elisabeth C. McGowan, MD; David P. Carlton, MD; Heidi M. Harmon, MD, MS; Jane E. Brumbaugh, MD; C. Michael Cotten, MD, MHS, Pablo J. Sánchez, MD; Anna Maria Hibbs, MD; Rosemary D. Higgins, MD; for the Eunice Kennedy Shriver National Institute of Child Health and Human Development Neonatal Research Network

at 5 Years in the OPTIMAL

ORIGINAL ARTICLE

JAMA | Original Investigation | CARING FOR THE CRITICALLY ILL PATIENT

Effect of Physiologic Point-of-Care Cardiopulmonary Resuscitation Training on Survival With Favorable Neurologic Outcome in Cardiac Arrest in Pediatric ICUs

A Randomized Clinical Trial

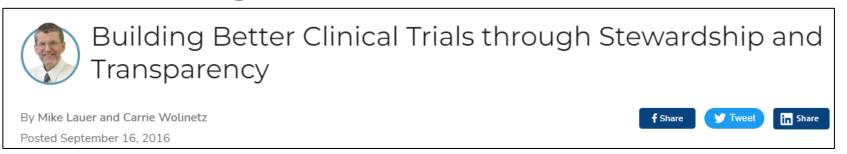
The ICU-RESUS and Eunice Kennedy Shriver National Institute of Child Health; and Human Development Collaborative Pediatric Critical Care Research Network Investigator Groups

etrozole versus Clomiphene for Infertility in the Polycystic Ovary Syndrome

ard S. Legro, M.D., Robert G. Brzyski, M.D., Ph.D., Michael P. Diamond, M.D., stos Coutifaris, M.D., Ph.D., William D. Schlaff, M.D., Peter Casson, M.D., ory M. Christman, M.D., Hao Huang, M.D., M.P.H., Qingshang Yan, Ph.D., uben Alvero, M.D., Daniel J. Haisenleder, Ph.D., Kurt T. Barnhart, M.D., right Bates, M.D., Rebecca Usadi, M.D., Scott Lucidi, M.D., Valerie Baker, M.D., russell, M.D., Stephen A. Krawetz, Ph.D., Peter Snyder, M.D., Dana Ohl, M.D., tte Santoro, M.D., Esther Eisenberg, M.D., M.P.H., and Heping Zhang, Ph.D., for the NICHD Reproductive Medicine Network*

Initiative Background





- NIH proposed to enhance clinical trial stewardship
- Individual Institutes/Centers asked to develop their own clinical trial FOAs
- Four principles for continued commitment of NICHD Network infrastructure for rigorous multisite clinical trials
 - Notice (NOT-HD-19-034): Infrastructure for NICHD Multisite Clinical Trials
 - https://grants.nih.gov/grants/guide/notice-files/NOT-HD-19-034.html
 - Webinar: Strategic Plan 2020 Implementation: NICHD's vision for multisite clinical trials infrastructure
 - https://www.youtube.com/watch?v=kzgFRuH9rnE



Infrastructure for NICHD Multisite Clinical Trials

Notice Number: NOT-HD-19-034

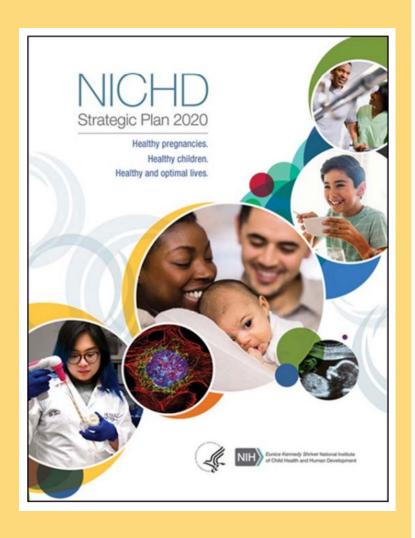
Key Dates

Release Date: October 10, 2019

Related Announcements

- •Enhance the rigor and reproducibility of clinical trial protocols
- •Promote greater availability of multisite clinical trial infrastructure to support trials from a wider range of investigators
- •Facilitate data sharing and access to biospecimens to efficiently expand research capacity for all investigators
- •Facilitate greater involvement of diverse populations in multisite clinical trials





NICHD Strategic Plan 2020

- Goal #1: Promoting an Inclusive Scientific Workforce
- Goal #2: Facilitating Data Sharing and Access to Biospecimens
- Goal #6: Improving Clinical Trial Oversight and Management



Initiative Background

NICHD

YEARS OF INNOVATION

- Process is needed to operationalize guiding principles
- NICHD conducted extensive assessment to inform this process
 - Request For Information (NOT-HD-19-041)
 - https://grants.nih.gov/grants/guide/notice-files/NOT-HD-19-041.html
 - Landscape Analysis of best clinical practices across the NIH
- Centralized approach
 - NICHD supports Network infrastructure
 - Any qualified investigator can use the infrastructure
 - All applications/protocols undergo NIH peer review
- A dedicated funding opportunity announcement is proposed to solicit clinical trial applications to leverage the NICHD Network Infrastructure





Enhancing rigor and reproducibility

- Develop new protocol proposal and selection processes to increase transparency and rigor of Network study selection
- Future trials to be conducted within the Network infrastructure:
 - Will be submitted as individual NIH applications
 - Will undergo NIH peer review
 - Funding decisions will be made by NICHD





Promoting greater availability of multisite clinical trial infrastructure to support trials from a wider range of investigators

- Investigators from outside the Network will now be able to propose studies to be conducted in the Network
- Investigators from Network Clinical Centers will also be eligible to propose studies via the proposed funding opportunity announcement
- Studies will need to align with the purpose, goals and objectives of the Network and NICHD priorities





Facilitating data sharing and access to biospecimens

- Networks will be required to share data and biospecimens according to the NIH Policy for Data Management and Sharing
- Study consent forms to allow broad sharing and future use of study data
- Common Data Element use will be a priority of Network studies



Facilitating greater involvement of diverse populations in multisite clinical trials

- Ensuring diverse populations in multisite clinical trials
 - Participants
 - Investigators
- Trials will be expected to maximize inclusion of diverse study participants
- Develop diversity in the workforce (<u>NOT-OD-20-031</u>)





- Essential features of that approach
 - High priority science aligned with our Networks
 - Rigorous pre-application process
 - Peer Review process
 - Funding procedure





- Scientific areas
 - Start with a small focus and expand progressively
 - Identify Network synergies
- Rigorous pre-application process
 - Feasibility assessment
 - Alignment of science
 - Current state of the field
 - Societal needs
 - Network mission
 - Institute priorities



NICHD Maternal-Fetal Medicine Units (MFMU) Network: Clinical Centers (UG1 Clinical Trial Optional)

UG1 Clinical Research Cooperative Agreements - Single Project

Reissue of RFA-HD-16-019

• May 19, 2022 - Notice of Pre-Application Webinars for the Maternal-Fetal Medicine Units Network and Neonatal Research Network Funding Opportunity Announcements. See Notice NOT-HD-22-023

RFA-HD-23-016

RFA-HD-23-017, U24 Resource-Related Research Project (Cooperative Agreements)

NICHD Neonatal Research Network (NRN): Data Coordinating Center (U24 Clinical Trial Optional)

U24 Resource-Related Research Projects – Cooperative Agreements

Reissue of RFA-HD-18-010

 May 19, 2022 - Notice of Pre-Application Webinars for the Maternal-Fetal Medicine Units Network and Neonatal Research Network Funding Opportunity Announcements. See Notice NOT-HD-22-023

RFA-HD-23-001

RFA-HD-23-002, UG1 Clinical Research Cooperative Agreements - Single Project

NICHD Neonatal Research Network (NRN): Clinical Centers (UG1 Clinical Trial Optional)

UG1 Clinical Research Cooperative Agreements - Single Project

Reissue of RFA-HD-16-020

 May 19, 2022 - Notice of Pre-Application Webinars for the Maternal-Fetal Medicine Units Network and Neonatal Research Network Funding Opportunity Announcements. See Notice NOT-HD-22-023

RFA-HD-23-002

RFA-HD-23-001, U24 Resource-Related Research Project (Cooperative Agreements)

NICHD Maternal-Fetal Medicine Units (MFMU) Network: Data Coordinating Center (U24 Clinical Trial Optional)

U24 Resource-Related Research Projects – Cooperative Agreements

Reissue of RFA-HD-13-013

 May 19, 2022 - Notice of Pre-Application Webinars for the Maternal-Fetal Medicine Units Network and Neonatal Research Network Funding Opportunity Announcements. See Notice NOT-HD-22-023

RFA-HD-23-017

RFA-HD-23-016, UG1 Clinical Research Cooperative Agreements - Single Project





- Scientific areas
 - Start with a small focus and expand progressively
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- Peer Review process
 - Network Clinical trials will undergo NIH peer review
 - Special emphasis panel
- Funding procedure
 - Cost neutral
 - Various models (e.g. StrokeNet)
 - Model that optimizes Network infrastructure



Moving forward



- Much work left to be done
- Expected Impact
 - Expand the breadth and diversity of our science and investigators
 - Enhance rigor and transparency of Network clinical trials
 - Perform clinical trials in the most efficient and cost-effective manner
 - Assure alignment with the NICHD Strategic Plan
 - Identify novel and innovative interventions
 - Foster nimbleness and flexibility for emerging health crises
 - Augment the impact of the science that we support



Team Effort



Core Group

- Abi Tepede
- Caroline Signore
- David Weinberg
- Yasaman Shirazi
- Dennis Twombly
- Bill Duval
- Kathryn Adams
- Rebekah Rasooly
- Rohan Hazra

Pre-Application Group

- Megan Mitchell
- Scott Janis

Funding Procedure

- Maggie Young
- Bonnie Jackson
- Kelly Fritz
- Mario Martinez
- Teri Pailen
- Alexis Clark
- Nichole Zivuku
- Melani Ledu

Scientific Content

- Nahida Chakhtoura
- Valerie Maholmes
- Tammara Jenkins
- Esther Eisenberg

Scientific Review

- Sherry Dupere
- Joanna Kubler-Kielb
- Cathy Wedeen

Clinical Trials Working Group

- Maria Stagnitto
- Sue Marden

Scientific Programmatic Branch Chiefs

Administrative Support

Ashley Walton





"It is not important for us to be right, but that we get it right."

- Rohan Hazra 2022



