

# THE LANCET **Neurology**

## **Supplementary webappendix**

This webappendix formed part of the original submission and has been peer reviewed. We post it as supplied by the authors.

Supplement to: Ginsberg MD, Palesch YY, Hill MD, et al, for the ALIAS and Neurological Emergencies Treatment Trials (NETT) Investigators. High-dose albumin treatment for acute ischaemic stroke (ALIAS) part 2: a randomised, double-blind, phase 3, placebo-controlled trial. *Lancet Neurol* 2013; published online Sept 27. [http://dx.doi.org/10.1016/S1474-4422\(13\)70223-0](http://dx.doi.org/10.1016/S1474-4422(13)70223-0).

## **Appendix**

### **Study Committees and Investigators**

#### **ALIAS Trial Executive Committee**

Myron D. Ginsberg, University of Miami, Miami, FL -- Study Chair and Principal Investigator, Clinical Coordinating Center;

Yuko Y. Palesch, Medical University of South Carolina, Charleston, SC -- Principal Investigator, Statistics and Data Coordinating Center;

Michael D. Hill, University of Calgary, Canada -- Director, Canadian Coordinating Center;

Bonnie D. Waldman, Medical University of South Carolina, Charleston, SC -- Project Manager;

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Richard Leinster, Medical University of South Carolina, Charleston, SC -- Data Manager;

Isabel Mendez, University of Miami, Miami, FL -- Financial Manager

Diego Tamariz, University of Miami, Miami, FL -- Clinical Project Coordinator;

Karla J. Ryckborst, University of Calgary, Calgary, Canada -- Study Coordinator;

Claudia S. Moy, NINDS, National Institutes of Health, Rockville, MD -- NINDS Project Scientist.

#### **Unblinded Study Statisticians**

Renee H. Martin, Medical University of South Carolina, Charleston, SC;

Sharon D. Yeatts, Medical University of South Carolina, Charleston, SC.

#### **External Safety Monitors**

Andrew M. Naidech, Northwestern University, Chicago;

Alejandro A. Rabinstein, Mayo Clinic, Rochester, MN.

#### **Data and Safety Monitoring Board**

Patrick D. Lyden, Cedars–Sinai Medical Center, Los Angeles, CA (Chair);

Christopher S. Coffey, University of Iowa, Iowa City, Iowa;

Marco Di Tullio, Columbia University, New York, NY;

Christine Wijman, Stanford University, Palo Alto, CA;

and Janice Cordell (ex officio), NINDS, National Institutes of Health, Rockville, MD.

## **PARTICIPATING CLINICAL SITES**

**[Site Principal Investigator, Primary Study Coordinator (number of subjects enrolled)]**

**[ \* denotes Neurological Emergency Treatment Trials (NETT) Network hub or spoke sites]**

### **UNITED STATES**

**Abington Memorial Hospital, Abington, PA\***: David Weisman, Jennifer McGowan Trainer (39);

**Memorial Hermann - Texas Medical Center, Houston, TX\***: Elizabeth Jones, Misty Ottman (36);

**University of Kentucky Medical Center, Lexington, KY\***: Roger Humphries, Linda Dechtenberg (33);

**Oregon Health and Science University, Portland, OR**: Wayne Clark, Darren Larsen (26);

**Mayo Clinic Hospital, Phoenix, AZ**: Maria Aguilar, Erica Boyd (24);

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**Metro Health Medical Center, Cleveland, OH:** Michael Bahntge, Scott Bailey (3);

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