



# KEMENTERIAN PENDIDIKAN NASIONAL UNIVERSITAS SEBELAS MARET

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## SURAT PENGANTAR

Nomor : 4994 /H27.5.2/PP/2010

Yth. Para Dekan Fakultas  
dilingkungan Universitas Sebelas Maret  
Surakarta

No.	Isi Surat / Hal	Banyaknya	Keterangan
1.	Copy surat dari Panitia International Seminar & Workshop UNDIP  Nomor : 08/ISW-FXA/VI/2010 Tanggal : 3 Juli 2010 Hal : Final Announcement	1 lembar	Dikirim dengan hormat, untuk diketahui dan mohon diinformasikan kepada staf pengajar yang berminat dilingkungan Fakultas Saudara.  Informasi lengkap melalui : <a href="http://www.cebjor.co.cc">www.cebjor.co.cc</a> , email ke <a href="mailto:Mrautism.UNDIP@gmail.com">Mrautism.UNDIP@gmail.com</a> atau telp. 0248412311  Terimakasih.

Surakarta, 14 JUL 2010

Kepala Bagian Kerjasama,

Drs. Pramista Magna  
NIP 196006071989031003

Tembusan Yth. :

1. Kepala International Office, UNS
2. Kabag. Sistem Informasi UNS  
(Mohon untuk dimuat di KONTAK dan website)

**2<sup>nd</sup> International Seminar & Workshop  
on Fragile-X, Autism and Related Disorders**



No. : 08/ISW-FXA/VI/2010  
Perihal : Final Announcement  
Lampiran : 1 berkas

03 JUL 2010

Kepada Yth,  
Rektor  
**Universitas Sebelas Maret**  
**Jl. Prof. Sutami 36 A**  
**Surakarta**

Diberitahukan dengan hormat, bahwa Universitas Diponegoro akan menyelenggarakan kegiatan *2nd International Seminar and Workshop on Fragile-X, Autism and Related Disorders*, yang dilaksanakan oleh Pusat Riset Biomedik (CEBIOR) FK UNDIP bekerjasama dengan *Medical Investigation of Neurodevelopmental Disorders (MIND) Institute, UC Davis*.

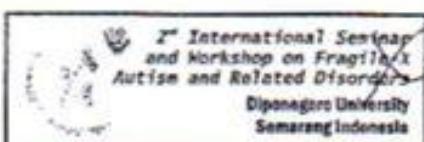
Adapun kegiatan tersebut akan diselenggarakan pada :

Hari/Tanggal : Sabtu, 7 Agustus 2010  
Waktu : 08.00 – 16.00 WIB  
Tempat : Patra Jasa Convention Hotel, Jl. Sisingamangaraja, Semarang

Mohon segera lakukan registrasi bila berminat. Kami juga mengundang seawat untuk mempresentasikan hasil riset yang berkaitan dengan Update dalam manajemen dan penanganan Retardasi Mental, Autism dan kelainan perkembangan saraf terkait melalui *Poster Presentation*. Batas pengumpulan abstrak hingga 7 Juli 2010. Info lengkap bisa diperoleh melalui: [www.cebiор.co.cc](http://www.cebiор.co.cc), email ke [MRautism.UNDIP@gmail.com](mailto:MRautism.UNDIP@gmail.com) atau telepon ke 024-8412311.

Terlampir adalah publikasi *Final announcement* untuk dapat disebarluaskan di lingkungan institusi Saudara. Atas perhatian dan kerjasama yang diberikan, kami ucapan terima kasih.

Hormat kami,  
Ketua Panitia



Prof. dr. Sultana MH Faradz, PhD  
NIP: 130 701 415

## Call for Poster

Theme : Update on management and treatment of MR, Autism, and other Neurodevelopmental Disorder

### Guidelines for submission of abstracts:

1. Abstracts may be submitted only via email : mrautism.undip@gmail.com
2. All abstracts must be submitted and written in English, maximum 250 words.
3. Poster size: 90 x 120 cm (portrait)
4. Abstracts must contain original material neither published nor presented elsewhere and must be approved by all contributing authors.
5. Please structure your abstract using the following headings: Introduction, Objectives, Methods, Results, Conclusion.
6. Abstracts will be subject to blind peer reviewers.
7. Deadline of abstract submission : July 7, 2010 and two weeks later, authors will be notified of acceptance by email.
8. Authors with accepted abstracts are entitled to register as participants.
9. Poster should be displayed by the participants in person one hour prior the registration and should be standby during the Poster Session.

## International Speakers

### Prof. Randi J. Hagerman, MD

Professor Randi Hagerman is a Developmental and Behavioral Pediatrician and the Medical Director of the M.I.N.D. Institute at UC Davis. She is internationally recognized as both a clinician and researcher in the Fragile X field. Professor Hagerman received her MD from Stanford University where she also carried out her Pediatric residency. She completed a Fellowship in Learning and Disabilities and Ambulatory

pediatrics at UC San Diego and, subsequently, spent the next 20 years from 1986 to 2006 at the University of Colorado where she headed Developmental and Behavioral Pediatrics. He co-founded the National Fragile X Foundation in 1984 in Colorado and developed a world-renowned Fragile X research and treatment center. In 2006, Professor Hagerman moved to UC Davis to be the Medical Director of the M.I.N.D. Institute. Dr. Hagerman and her team discovered the Fragile X-associated Tremor Ataxia Syndrome (FXTAS) which is a neurological disorder that affects older carriers of fragile X.

### Prof. Paul J. Hagerman, MD, PhD

He received both his M.D. and Ph.D. degrees from Stanford University School of Medicine in 1977. This was followed by a three year Leukemia Society Fellowship at UCSF, where he began his studies of nucleic acid structure. He joined the University of Colorado in 1980, where he remained prior to joining the UC Davis faculty. Hagerman is currently on the scientific advisory board of the Muscular Dystrophy Association, and recently started toward the study of the

molecular biology of neurodevelopmental disorders, with particular emphasis on fragile X syndrome, the leading heritable form of mental retardation. Since joining the Davis faculty, Dr. Hagerman has expanded his research efforts to include investigations of the underlying biochemical and genetic causes of autism.

### Louise Gane, M.Sc.

She is Senior Genetic Associate at the M.I.N.D. Institute. In 1991, she joined the team led by Dr. Hagerman at the Fragile X Research and Treatment Center in Denver, Colorado as a genetic counsellor. Under the direction of Dr. Randi Hagerman, Louise is now located at the UC Davis M.I.N.D. Institute working as a Genetic Counsellor and Assistant to the Director. She is involved in counselling patients with Fragile X syndrome and their family members. In her role Louise helps families deal with and understand the diagnosis, assessing genetic risk, and explaining reproductive options.

## Committee

### Patron

Rector of Diponegoro University

Director of Dr Kariadi Hospital Semarang

Dean, Faculty of Medicine Diponegoro University (FMUDU)

Vice Rector for Development and Cooperation

Vice Dean for Finance

Advisor of Center for Biomedical Research (CeBior)

### Chair

Prof. Sulfana MH Farez, MD, PhD

### Co-Chair

Fitri Hartanto, MD (Pediatrician)

### Secretary

Neni Sustiseninggih, MD, Msc

Mulihulah Muniroh, MD, MSc

### Secretariat

Nani Mahatani, MD, Msc, Ardina Apriiani, S.Sos

### Treasurer

Ratna Damma Purnawati, MD, Msc, Annastasia Edlati, S.Psi, MSc, Psi

### Scientific Committee

Astri Purwanti, MD (Pediatrician), Aifiani Hikmah Putranti, MD (Pediatrician), Dani Rahmawati, MD (Neurologist), Afifati Fitriksasari, MD (Psychiatrist), Hexantio, MD (Neurologist), Eko Adhi Pangarsa, MD (Internist), Yetti Movieta Nenicy, MD (Pediatrician), Farmadyta EP Mundofia, MD, Msc, Maria Belladona, MD, M. Meritalia Setiawati, MD, PhD (Pediatrician), Rudi Handoyo, MD (Medical rehabilitation)

### Talk show Committee

Agustini Utami, MD, MSc (Pediatrician), Costrie Ganes, SPsi

Audiovisual & Documentation

Ahmad Zulfa Junianto, MD, MSc (Andrologist)

### Accommodation

Aris Catur, MD (Neurologist)

### Translator

Widagdo, S.Ked

# 2<sup>nd</sup> International Seminar at Workshop on Fragile-X, Autism and Related Disorders



Semarang, 7<sup>th</sup> August 2010

Diponegoro University

Medical Investigation of  
Neurodevelopmental Disorders  
(MIND) Institute, UC Davis



Widagdo, S.Ked

## Background

Fragile-X syndrome (FXS) is the most common form inherited intellectual disabilities (ID). The clinical signs include intellectual disabilities, hyperactive, autism, psychoneurologic disorders, and macroorchidism (testes enlargement) in male. The common affected is in males; however, carrier female could have a mild intellectual disability such as learning difficulties, without other symptoms. Some clinically normal males (Normal transmitting Males) are known to carry and transmit the fragile X mutation to their daughter. In advance age, carriers of FXS could develop Fragile-X Associated Tremor Ataxia (AS), a neurodegenerative disorder.

Autistic spectrum disorder (ASD) involves developmental abnormalities in reciprocal social interaction, communication, and repetitive behavior. Some genetic disorders associated with autism for example: in Syndrome (DS), FXS and microdeletion chromosomal disorders such as Angelman syndrome, Prader Willi syndrome etc. The information gap between experts and patients could be a serious problem for patient management. Recently many studies in this field have been developed to find the newest and effective treatment and management.

Advances in understanding the neurobiology of FXS have led to new targeted treatments. The most remarkable is the metabotropic glutamate receptor 5 (mGluR5) antagonists. The mGluR5 antagonists have been studied in animal models of fragile X and have been shown to have a beneficial effect on seizures, cognition, and behavior. Other recent advances in FXS treatment is Minocycline. Studies in animal models suggest that minocycline is not useful as an antibiotic but may also has potential as a protective agent. Those advancements in FXS treatment in patients will be discussed in this workshop.

Within the framework of long collaboration with the Medical Investigation of Neurodevelopmental Disorders (MIND) Institute, UC Davis, USA; Faculty of Medicine, Oregon University (FMDU) Semarang Indonesia will organize a 2nd International workshop on Fragile X syndrome, Autism and Related Disorders. This workshop

comprises also a talk show session, which will present some cases of autism and ID patients.

## Purpose

To disseminate recent advances and treatments in the field of Fragile X syndrome, Autism and related disorders from the expert scientist to the parents and people who are interested in those field.

## Venue

Patra Convention Hotel  
Jl. Sisingamangaraja, Semarang

## Participants

This workshop is suited for parents, students (undergraduates and postgraduates), medical scientists, clinicians, nutritionist, psychologist, and people who are interested. The number of participants is limited to 300 people.

## Date

August, 7th 2010

## Registration Fee

UNDIP staff	Free
Non UNDIP staff	
Physician (Specialist)	Rp. 500.000,-
General Practitioner	Rp. 250.000,-
Public	Rp. 250.000,-
Discount 50% for autism parents, FMDU students and alumni of genetic counselling master program UNDIP.	
Information and registration	
Wiwik / Dina (Center for Biomedical Research) GSG 2nd Floor Jl. Dr. Sutomo 14 Semarang Tel: +62-24-8412311 Fax: +62-24-8454714 HP:+62-81-931980014 e-mail: molcytogen@yahoo.com, genetikaundip@indosat.net.id website : www.cebjor.co.cc	

## Program

Saturday, August 7, 2010	
07.30 - 08.00	Registration
08.00 - 08.15	Opening Speech by: 1. Chairperson 2. Rector of Diponegoro University
08.15 - 08.30	Future and challenges of Autism Center Indonesia Prof. Susilo Wibawa, MD, PhD, Androlog
08.30 - 08.50	Fragile X syndrome in Indonesia Prof. Sultana MH Faradz, MD, PhD
08.50 - 09.20	RNA toxicity and FXTAS in Fragile-X premutation carriers Prof. Paul Hagerman, MD, PhD
09.20 - 09.50	Clinical Aspect of Autism and Its Management Melly Budhirman, MD, Child Psychiatrist
09.50 - 10.20	Autism and Fragile X syndrome: Advances in Targeted treatments Prof. Randi J. Hagerman, MD, PhD
10.20 - 10.35	Discussion
10.30 - 11.05	Minocycline study in Fragile X syndrome and Autism Agustini Utari, MD, M.Sc, Pediatrician Psychopharmacological Intervention in Autism and Fragile X syndrome Prof. Randi J. Hagerman, MD, PhD Genetic Counseling in Fragile X syndrome and Autism
11.05 - 11.45	Louise Game, M.Sc Discussion
12.25 - 13.10	Lunch and poster session
13.10 - 15.20	Talk show (patients, parents and experts)
15.20 - 15.40	Best posters announcement and Closing ceremony