

CTAD 2012

Clinical Trials on Alzheimer's Disease

Montpellier '08 | Las Vegas '09 | Toulouse '10 | San Diego '11 | Monte Carlo '12

Clinical Trials on Alzheimer's Disease

October 29-31, 2012
Forum Grimaldi - Monte Carlo

CME CREDITS

Clinical Trials



**CTAD
Monte Carlo
2012**



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MONTPELLIER and TOULOUSE EADC Centers
European Alzheimer's Disease Consortium



UC San Diego
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Dear Colleague,

We are proud to present our program for this 5th edition of Clinical Trials on Alzheimer's Disease which will take place October 29-31, 2012 at the Grimaldi Forum in Monte Carlo with the support of the Monegasque Association for research on Alzheimer's disease (AMPA)

Alzheimer's disease is one of the most important health challenges facing aging populations worldwide. The development of the next generation of Alzheimer's disease drugs is becoming essential to face up to this challenge. New pathways have been identified with biomarkers, facilitating novel trial designs for studies of tau-based therapies and other disease-modifying drugs including immunotherapy. However, methodological challenges continue to slow the development of specific new drug candidates. One of the objectives of the conference is to identify these hurdles and find ways to address them by bringing together world leaders in AD drug development to discuss solutions to the difficulties that have slowed the pace of progress, with a particular focus on clinical trial methodology.

Again this year CTAD is the perfect opportunity to hear about the clinical experiences of international teams, exchange with your peers on the difficulties and challenges of Alzheimer's disease and take home some hands-on therapeutic and methodological tools to improve and reinforce your AD clinical trials teams. CME Credit are available from the University of California at San Diego (USCD).

We are very happy to welcome you to CTAD 2012!

Paul Aisen - Jacques Touchon - Bruno Vellas - Michael Weiner

The Monegasque Association for research on Alzheimer's disease (AMPA) is a non-profit organization chaired by Catherine Pastor and Vice-President Prof. Alain Pesce, head of the Department of Geriatrics at the Princess Grace Hospital in Monaco. AMPA's mission is to support research on Alzheimer's disease, to sustain local initiatives for people with dementia and their caregivers, to raise awareness and disseminate information and expertise. Since 2008, AMPA also promotes international exchanges between scientists in order to increase knowledge and encourage innovative research.

www.ampa-monaco.com

Clinical Trials on Alzheimer's Disease

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ORGANIZING COMMITTEE

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Monday, October 29th

8.00 - 8.15 a.m	WELCOME BY THE ORGANIZING COMMITTEE
8.15 - 8.45 a.m	KEYNOTE 1 : 5-year experience in AD clinical trials as member of the Medical Committee for Medical Committee for Medicinal Products for Human Use and CNS working group within the EMEA
8.45 - 9.45 a.m	Symposium 1 : Harmonizing Regulatory Requirements to Benefit Future Alzheimer's Disease Patients
9.45 - 10.45 a.m	ORAL COMMUNICATIONS <i>Coffee Break and poster sessions</i>
11.15 - 12.15 p.m	Symposium 2 : Bapineuzumab IV Phase 3 Results <i>Lunch Break and poster sessions</i>
1.30 - 2.30 p.m	Symposium 3 : Effects of Apolipoprotein E Isoforms on Patient Characteristics and Trial Outcomes in Light of Recent Phase 3 results : Stratified Medicine for Alzheimer's Disease Drug Development
2.30 - 4.00 p.m	ORAL COMMUNICATIONS <i>Coffee Break and poster sessions</i>
4.30 - 6.10 p.m	UPDATE ON CLINICAL TRIALS 1
7.00 p.m	WELCOME RECEPTION with the support of AMPA

Tuesday, October 30th

8.00 - 8.45 a.m	ORAL COMMUNICATIONS
8.45 - 9.45 a.m	Symposium 4 : Enrichment Approaches in Predementia AD Clinical Trials
9.45 - 10.25 a.m	UPDATE ON CLINICAL TRIALS 2
	<i>Coffee Break and poster sessions</i>
11.00 - 12.00 p.m	Symposium 5 : Solanezumab Phase 3 results: Implications for Alzheimer's disease modification
	<i>Lunch Break and poster sessions</i>
1.00 - 3.00 p.m	ORAL COMMUNICATIONS
3.00 - 4.00 p.m	Symposium 6 : Modeling the Course of AD : Contributions to Better Clinical Trials
	<i>Coffee Break and poster sessions</i>
4.30 - 5.30 p.m	Symposium 7 : MAPT (Multidomain Alzheimer Preventive Trial) imaging data
5.30 - 6.00 p.m	KEYNOTE 2 : New design for symptomatic treatment of AD
6.00 - 6.30 p.m	ORAL COMMUNICATIONS

Wednesday, October 31st

8.00 - 9.00 a.m	UPDATE ON CLINICAL TRIALS 3
9.00 - 10.00 a.m	ORAL COMMUNICATIONS
	<i>Coffee Break and poster sessions</i>
10.30 - 12.30 p.m	ORAL COMMUNICATIONS

Monday, October 29th

8.00 - 8.15 a.m	<p>WELCOME BY THE ORGANIZING COMMITTEE <i>Paul Aisen, University of California at San Diego, USA, Jacques Touchon, Montpellier University Hospital, INSERM U1061, France, Bruno Vellas, Toulouse University Hospital, France, Mike Weiner, University of California at San Francisco, USA</i></p>
8.15 - 8.45 a.m	<p>KEYNOTE 1 : 5-year experience in AD clinical trials as member of the Medical Committee for Medical Committee for Medicinal Products for Human Use and CNS working group within the EMEA <i>Cristina Sampaio, Scientific Expert, EMEA</i></p>
8.45 - 9.45 a.m	<p>Symposium 1 : Harmonizing Regulatory Requirements to Benefit Future Alzheimer's Disease Patients Chairman : Daniel Perry ⁽¹⁾</p> <p>Presenters : Diane Stephenson ⁽²⁾, Karl Broich ⁽³⁾ ⁽¹⁾ ACT-AD Coalition, Alliance for Aging Research, USA, ⁽²⁾ Coalition Against Major Diseases, Critical Path Institute, USA ⁽³⁾ Federal Institute for Drugs and Medical Devices (BfArM)</p>
9.45 - 10.45 a.m	<p>ORAL COMMUNICATIONS Moderators : M.Egan, H.Hampel</p>
9.45 - 10.00 a.m	<p>OC1 - Is the Dementia of Alzheimer's Disease due to the Toxicity of β-Amyloid or Tau? The Implications of this Question for Drug Discovery <i>Jordan Holtzman, University of Minnesota, Minneapolis, MN, USA</i></p>
10.00 - 10.15 a.m	<p>OC2 - Potential of Human analogue of Morris Water Maze in translational medicine and the assessment of therapeutic response <i>John Harrison ^(2,4,5), Jan Laczó ^(1,2), Manfred Windisch ^(2,3), Jakub Hort ^(1,2)</i> ⁽¹⁾ Memory Disorders Clinic, Department of Neurology, Charles University in Prague, 2nd Medical Faculty and University Hospital Motol, Prague, Czech Republic, ⁽²⁾ Polyhymnia Translational Research, London, UK, ⁽³⁾ JSW-Lifesciences GmbH, Grambach-Graz, Austria, ⁽⁴⁾ Metis Cognition Ltd., Kilmington, UK, ⁽⁵⁾ Dept. of Medicine, Imperial College, London, UK</p>
10.15 - 10.30 a.m	<p>OC3 - Assumptions of mortality have a great impact on the cost-effectiveness of disease-modifying drugs in AD <i>Anders Sköldunger ⁽¹⁾, Kristina Johnell ⁽¹⁾, Bengt Winblad ⁽²⁾, Anders Wimo ⁽²⁾</i> ⁽¹⁾ Aging Research Center, Karolinska Institutet and Stockholm University, Stockholm, Sweden, ⁽²⁾ KI-Alzheimer's Disease Research Center, Karolinska Institutet, Huddinge, Sweden</p>

Monday, October 29th *Continued*

10.30 - 10.45 a.m	<p>OC4 - The Novel BACE Inhibitor MK-8931 Dramatically Lowers CSF Aβ Peptides in Healthy Subjects Following Single and Multiple Dose Administration <u>Michael F. Egan</u>, Mark S Forman ⁽¹⁾, John Palcza ⁽¹⁾, Jack Tseng ⁽¹⁾, Jos Leempoels ⁽²⁾, Steven Ramael ⁽²⁾, David Han ^(3,4), Stanford Jhee ⁽³⁾, Larry Ereshefsky ^(3,5), Michael Tanen ⁽¹⁾, Omar Laterza ⁽¹⁾, Marissa Dockendorf ⁽¹⁾, Gopal Krishna ⁽¹⁾, Lei Ma ⁽¹⁾, John A Wagner ⁽¹⁾, Matthew D Troyer ⁽¹⁾ ⁽¹⁾ Merck, Whitehouse Station, NJ, USA, ⁽²⁾ SGS Life Science Services, Antwerpen, Belgium, ⁽³⁾ Parexel International Early Phase, Glendale, CA, USA, ⁽⁴⁾ California Clinical Trials Medical Group, Glendale, CA, ⁽⁵⁾ University of Texas Health Science Center, San Antonio, TX</p>
10.45 - 11.15 a.m	Coffee Break and poster sessions
11.15 - 12.15 p.m	<p>Symposium 2 : Bapineuzumab IV Phase 3 results Chairman : Philip Scheltens ⁽¹⁾</p>
	<p>Presenters : Reisa Sperling ⁽²⁾, Stephen Salloway ⁽³⁾, Nick Fox ⁽⁴⁾ ⁽¹⁾ VU University Medical Center, Alzheimer Center, Amsterdam, the Netherlands, ⁽²⁾ Brigham & Women's Hospital, Boston, MA, USA ⁽³⁾ Butler Hospital, Providence, RI, USA, ⁽⁴⁾ UCL, Institute of Neurology, London, United Kingdom</p>
12.15 - 1.30 p.m	Lunch Break and poster sessions
1.30 - 2.30 p.m	<p>Symposium 3 : Effects of Apolipoprotein E Isoforms on Patient Characteristics and Trials Outcomes in Light of Recent Phase 3 Results : Stratified Medicine for Alzheimer's Disease Drug Development Chairman : Lon.S.Schneider, University of Southern California, Los Angeles, CA, USA</p>
1.30 - 1.50 p.m	<ul style="list-style-type: none"> The neurobiology and impact of apoE4 and apoE2 carriage on clinical trials Terry Goldberg, Hofstra North Shore LIJ School of Medicine, Manhasset, New York, USA
1.50 - 2.10 p.m	<ul style="list-style-type: none"> Relation of apoE to brain structure and function in Alzheimer's disease and aging Michela Pievani, IRCCS Fatebenefratelli Brescia, Brescia, Italy
2.10 - 2.30 p.m	<ul style="list-style-type: none"> Simulating apoE stratified medicine trials in Alzheimer's disease Lon S.Schneider ⁽¹⁾, Richard Kennedy ⁽²⁾, Gary Cutter ⁽²⁾ ⁽¹⁾ Keck School of Medicine of the University of Southern California, Los Angeles, CA, USA, ⁽²⁾ School of Public Health, University of Alabama, Birmingham, Alabama, USA

Monday, October 29th *Continued*

2.30 - 4.00 p.m

ORAL COMMUNICATIONS

Moderators : *G.Frisoni, B.Winblad*

2.30 - 2.45 p.m

OC5 - Measuring cognitive change from mild cognitive impairment to prodromal Alzheimer disease

T.Mura ^(1,2,3,4), C.Proust-Lima ^(5,6), H.Jacqmin-Gadda ^(5,6), TN.Akbaraly ^(1,2,7), B.Dubois ⁽⁸⁾, C.Berr ^(1,2,9)
⁽¹⁾ INSERM, U1061, Neuropsychiatrie : Recherche Epidémiologique et Clinique, Montpellier, France, ⁽²⁾ Université Montpellier I, Montpellier, France, ⁽³⁾ Département d'Information Médicale, Centre d'Investigation Clinique, CHU Montpellier, Montpellier, France, ⁽⁴⁾ INSERM, CIC 1001, Montpellier, France, ⁽⁵⁾ INSERM U897, Equipe de Biostatistique, Centre de Recherche en Epidémiologie et Biostatistique, Bordeaux, France, ⁽⁶⁾ Université Bordeaux Segalen, ISPED, Bordeaux, France, ⁽⁷⁾ Department of Epidemiology and Public Health, University College London, London, United Kingdom, ⁽⁸⁾ INSERM-UPMC UMRS 975, Institut de la Mémoire et de la Maladie d'Alzheimer, ICM, APHP, Salpêtrière Hospital, Université Paris 6, Paris, France, ⁽⁹⁾ CMRR Languedoc Roussillon, service de Neurologie, CHU Montpellier, Montpellier, France

2.45 - 3.00 p.m

OC6 - Long-Term Longitudinal Biomarker Trials in Subjects at Genetic Risk of Developing Alzheimer's Disease : the GEPARD-AD Studies

Joel Ross ⁽¹⁾, Paul M.Thompson ⁽²⁾, Rachelle S.Doody ⁽³⁾, Pierre N.Tariot ⁽⁴⁾, Eric M.Reiman ⁽⁴⁾, Jessica Langbaum ⁽⁴⁾, Lon S.Schneider ⁽⁵⁾, Ugo Lucca ⁽⁶⁾, Enrico Frigerio ⁽⁷⁾, Francesco Fiorentini ⁽⁷⁾, Luciana Giardino ⁽⁸⁾, Laura Calzà ⁽⁸⁾, Dottie Norris ⁽⁹⁾, Helen Cicirello ⁽⁹⁾, Daniela Casula ⁽⁹⁾, Bruno P.Imbimbo ⁽⁹⁾
⁽¹⁾ Memory Enhancement Center of America, 4 Industrial Way West, Eatontown, NJ, USA, ⁽²⁾ Laboratory of Neuro Imaging, Dept. of Neurology, UCLA School of Medicine, Los Angeles, CA, USA, ⁽³⁾ Alzheimer's Disease and Memory Disorders Center, Baylor College of Medicine, Department of Neurology, Houston, TX, USA, ⁽⁴⁾ Banner Alzheimer's Institute, AZ, USA, ⁽⁵⁾ University of Southern California Keck School of Medicine, Los Angeles, CA, USA, ⁽⁶⁾ Istituto di Ricerche Farmacologiche Mario Negri, Milano, Italy
⁽⁷⁾ Accelera Srl, Nerviano, Italy, ⁽⁸⁾ Health Sciences and Technologies - Interdepartmental Center for Industrial Research (HST-ICIR), University of Bologna, Ozzano Emilia, Bologna, Italy, ⁽⁹⁾ Research & Development, Chiesi Pharmaceuticals Inc., Rockville, MD 20850, USA

3.00 - 3.15 p.m

OC7 - The Dominantly Inherited Alzheimer's Network Trials

Randall Bateman, on behalf of the Dominantly Inherited Alzheimer Network

3.15 - 3.30 p.m

OC8 - Definition of Harmonized Protocol for Hippocampal Segmentation

Marina Boccardi ⁽¹⁾, Martina Bocchetta ^(1,2), Liana Apostolova ⁽³⁾, Josephine Barnes ⁽⁴⁾, George Bartzokis ⁽⁵⁾, Gabriele Corbetta ⁽¹⁾, Charles DeCarli ⁽⁶⁾, Leyla deToledo-Morrell ⁽⁷⁾, Michael Firbank ⁽⁸⁾, Rossana Ganzola ⁽¹⁾, Lotte Gerritsen ⁽⁹⁾, Wouter Henneman ⁽¹⁰⁾, Ronald J.Killiany ⁽¹¹⁾, Nikolai Malykhin ⁽¹²⁾, Patrizio Pasqualetti ⁽²⁾, Jens C.Pruessner ⁽¹³⁾, Alberto Redolfi ⁽¹⁾, Nicolas Robitaille ⁽¹⁴⁾, Hilikka Soininen ⁽¹⁵⁾, Daniele Tolomeo ⁽¹⁾, Lei Wang ⁽¹⁶⁾, Craig Watson ⁽¹⁷⁾, Henrike Wolf ⁽¹⁸⁾, Simon Duchesne ⁽¹⁴⁾, Clifford R.Jack Jr ⁽¹⁹⁾, Giovanni B.Frisoni ⁽¹⁾
⁽¹⁾ LENITEM (Laboratory of Epidemiology, Neuroimaging and Telemedicine) IRCCS – S. Giovanni di Dio – Fatebenefratelli Brescia, Italy, ⁽²⁾ AfAR – Associazione Fatebenefratelli per la Ricerca, Rome, Italy, ⁽³⁾ Laboratory of Neuroimaging, David Geffen School of Medicine, University of California, Los Angeles, CA, ⁽⁴⁾ Dementia Research Centre, UCL Institute of Neurology, University College London, London, UK, ⁽⁵⁾ Department of Psychiatry, David Geffen School of Medicine at UCLA, Los Angeles, CA, ⁽⁶⁾ Department of Neurology, University of California, Davis, CA, ⁽⁷⁾ Department of Neurological Sciences, Rush University, Chicago, Illinois, ⁽⁸⁾ Institute for Ageing and Health, Newcastle University, Wolfson Research Centre, Newcastle, UK, ⁽⁹⁾ Karolinska Institute, Stockholm, Sweden, ⁽¹⁰⁾ Department of Radiology and Alzheimer Center, VU University Medical Center, Amsterdam, The Netherlands, ⁽¹¹⁾ Department of Anatomy and Neurobiology, Boston University School of

Monday, October 29th *Continued*

2.30 - 4.00 p.m	ORAL COMMUNICATIONS <i>Continued</i>
	<p><i>Medicine, ^(1,2) Department of Biomedical Engineering, Centre for Neuroscience, University of Alberta, Edmonton, Alberta, Canada, ⁽¹³⁾ McGill Centre for Studies in Aging, Department of Psychiatry, McGill University, Montreal, Quebec, Canada, ⁽¹⁴⁾ Department of Radiology, Université Laval and Centre de Recherche Université Laval – Robert Giffard, Quebec City, Canada, ⁽¹⁵⁾ Dept of Neurology, University of Eastern Finland and Kuopio University Hospital, Kuopio, Finland, ⁽¹⁶⁾ Northwestern University Feinberg School of Medicine, Department of Psychiatry and Behavioral Sciences, United States, ⁽¹⁷⁾ Wayne State University School of Medicine, D-University Health Center, St. Antoine, Detroit, MI, ⁽¹⁸⁾ Department of Psychiatry Research and Geriatric Psychiatry, Psychiatric University Hospitals, University of Zurich, Zurich, Switzerland, ⁽¹⁹⁾ Department of Diagnostic Radiology, Mayo Clinic and Foundation, Rochester, MN</i></p> <p>3.30 - 3.45 p.m OC9 - Robustness of Automated Hippocampal Volumetry across MR Field strengths and Same-Day Repeat Scans <u>Robin Wolz</u> ^(1,2), Adam J.Schwarz ⁽³⁾, Peng Yu ⁽³⁾, Pat Cole ⁽⁴⁾, Daniel Rueckert ^(1,2), David Raunig ⁽⁵⁾, Derek Hill ⁽¹⁾ ⁽¹⁾ IXICO Ltd, London, UK, ⁽²⁾ Imperial College London, London, UK, ⁽³⁾ Elli Lilly and Company, Indianapolis IN, USA, ⁽⁴⁾ Cole Consulting, NJ, USA, ⁽⁵⁾ Icon Medical Imaging, Warrington PA, USA</p> <p>3.45 - 4.00 p.m OC10 - Disentangling the normal aging from the pathological Alzheimer's disease progression on structural MR images <u>Marco Lorenzi</u> ^(1,2), Nicholas Ayache ⁽²⁾, Xavier Pennec ⁽²⁾, Giovanni B.Frisoni ⁽¹⁾, and the Alzheimer's Disease Neuroimaging Initiative (ADNI) ⁽¹⁾ IRCCS San Giovanni di Dio Fatebnefratelli, Brescia, Italy, ⁽²⁾ Asclepios research project, INRIA Sophia antipolis, France</p>
4.00 - 4.30 p.m	<i>Coffee Break and poster sessions</i>
4.30 - 6.10 p.m	UPDATE ON CLINICAL TRIALS 1 Moderators : R.Doody, J.Touchon
4.30 - 4.50 p.m	1 - Safety and Efficacy of Solanezumab in Patients with Mild to Moderate Alzheimer's Disease: Results from Phase 3 <i>Rachelle S.Doody, Baylor College of Medicine - Department of Neurology, Houston, Texas, USA</i>
4.50 - 5.10 p.m	2 - The Effects of ELND005 (Scyllo-inositol) on Agitation/Aggression in AD Dementia : Results from a 78 week Phase 2 Study in Mild to Moderate AD <u>Susan Abushakra</u> ⁽¹⁾ , Constantine Lyketsos ⁽²⁾ , Gerald Crans ⁽¹⁾ , Priya Nadkarni ⁽¹⁾ , Chito Hernandez ⁽¹⁾ , Bruno Vellas ⁽³⁾ ⁽¹⁾ Elan Pharmaceuticals, Inc., South San Francisco, CA, USA, ⁽²⁾ Johns Hopkins University, Baltimore, MD, USA, ⁽³⁾ University of Toulouse, Toulouse, France
5.10 - 5.30 p.m	3 - ERP as biomarkers for Preclinical-AD <i>Karim Bennys ⁽¹⁾, Audrey Gabelle ⁽¹⁾, Jacques Touchon ^(1,2), ⁽¹⁾ Department of Neurology, Memory Research Resource Center for Alzheimer's Disease, University Hospital of Montpellier, France, ⁽²⁾ Inserm U1061, Montpellier, France</i>

Monday, October 29th *Continued*

4.30 - 6.10 p.m	UPDATE ON CLINICAL TRIALS <i>Continued</i>
5.30 - 5.50 p.m	4 - The SKT short cognitive performance test in pre-clinical and dementia stages of neurocognitive disorders <i>Mark Stemmler, University of Erlangen-Nuremberg, Germany</i>
5.50 - 6.10 p.m	5 - Applicability of new criteria for Clinical Trials in AD <i>Bruno Dubois⁽¹⁾, Paul Aisen⁽²⁾</i> <i>⁽¹⁾ Dementia Research Center, Salpêtrière University Hospital, Paris, France, ⁽²⁾ Department of Neurosciences, UCSD, San Diego, USA</i>
7.00 p.m	Bus departure for the WELCOME RECEPTION with support of The Monegasque Association for research on Alzheimer's disease (AMPA) - <i>Catherine PASTOR</i>

Tuesday, October 30th

8.00 - 8.45 a.m	ORAL COMMUNICATIONS Moderators : <i>S.Hendrix, P.Scheltens</i>
8.00 - 8.15 a.m	OC11 - Bayesian adaptive trial design: A new approach for Phase 2 clinical trials in Alzheimer's disease <i>Andrew Satlin⁽¹⁾, Veronika Logovinsky⁽¹⁾, Jinping Wang⁽¹⁾, Chad Swanson⁽¹⁾, Scott Berry⁽²⁾, Don Berry⁽²⁾</i> <i>⁽¹⁾ Eisai Inc., Neuroscience Product Creation Unit, NJ, ⁽²⁾ Berry Consultants, LLC, Austin, TX USA</i>
8.15 - 8.30 a.m	OC12 - A New Tool for Optimizing Responsiveness to Decline in Early AD <i>Suzanne Hendrix⁽¹⁾, Veronika Logovinsky⁽²⁾, Carlos Perdomo⁽²⁾, Jinping Wang⁽²⁾, Andrew Satlin⁽²⁾</i> <i>⁽¹⁾ Pentara Corporation, Salt Lake City, UT, 84109, ⁽²⁾ Eisai Inc., Neuroscience Product Creation Unit, Woodcliff Lake, NJ</i>
8.30 - 8.45 a.m	OC13 - Effects of age and APOE ε4 on the course of clinical and structural decline in Alzheimer disease <i>Dominic Holland⁽¹⁾, Linda K McEvoy⁽²⁾, Rahul S.Desikan⁽²⁾, Anders M.Dale^(1,2)</i> <i>⁽¹⁾ Department of Neurosciences, ⁽²⁾ Department of Radiology, University of California, San Diego, CA, USA</i>

Tuesday, October 30th

8.45 - 9.45 a.m	Symposium 4 : Enrichment Approaches in Predementia AD Clinical Trials Chairman : Jesse Cedarbaum ⁽¹⁾
8.55 - 9.20 a.m	<ul style="list-style-type: none"> • Predementia AD: Qualifying CSF and PET biomarkers for clinical trial use <i>Remy Cahn ⁽³⁾</i>
9.25 - 9.45 a.m	<ul style="list-style-type: none"> • Experience with CSF and PET Biomarkers for Enrichment of Predementia AD Trials <i>Christopher van Dyck ⁽³⁾</i> <p><small>⁽¹⁾ Global Clinical Research Bristol Meyers Squibb, Princeton, NJ, USA, ⁽²⁾ Global Regulatory Science, Bristol-Myers Squibb, Rueil, France, ⁽³⁾ Yale University, New Haven, CT, USA</small></p>
9.45 - 10.25 a.m	UPDATE ON CLINICAL TRIALS 2 Moderators : E. Giacobini, L. Buée
9.45 - 10.05 a.m	6 - Harmonisation of reporting standards for studies of diagnostic test accuracy in dementia : the STARDdem (STAndards for the Reporting of Diagnostic accuracy studies-Dementia) criteria Description of STARDdem process and Validation Study <u>Rupert McShane</u> ⁽¹⁾, Anna Noel-Storr ⁽¹⁾, Leon Flicker ⁽²⁾, Craig W Ritchie ⁽³⁾, Gordon Wilcock ⁽⁴⁾, STARDdem Working Group <small>⁽¹⁾ Cochrane Dementia and Cognitive Improvement Group, University of Oxford, United Kingdom, ⁽²⁾ Geriatric Medicine, University of Western Australia ⁽³⁾ Old age psychiatry, Imperial College, London, ⁽⁴⁾ Clinical Geratology, University of Oxford</small>
10.05 - 10.25 a.m	7 - Is the Beta-Amyloid cascade hypothesis dead ? If so, can it be resuscitated ? Ezio Giacobini, University of Geneva, Faculty of Medicine, Department of geriatrics
10.25 - 11.00 a.m	<i>Coffee Break and poster sessions</i>
11.00 - 12.00 p.m	Symposium 5 : Solanezumab Phase 3 results: Implications for Alzheimer's disease modification Chairman : Eric Siemers, Eli Lilly, USA
Noon to 1 p.m	<i>Lunch Break and poster sessions</i>

Tuesday, October 30th

1.00 - 3.00 p.m	ORAL COMMUNICATIONS Moderators : J.Hugon, D.Holland	
1.00 - 1.15 p.m	OC14 -	Enrichment, Stratification, and Outcome Measures for Predementia Alzheimer Disease Clinical Trials <u>Dominic Holland</u> ⁽¹⁾ , Linda K McEvoy ⁽²⁾ , Rahul S.Desikan ⁽²⁾ , Anders M.Dale ^(1,2) <i>⁽¹⁾ Department of Neurosciences, ⁽²⁾ Department of Radiology, University of California, San Diego, CA, USA</i>
1.15 - 1.30 p.m	OC15 -	Missing data in ADCS clinical trials <u>Michael Donahue</u> ^(1,2) , Anthony Gamst ^(1,2) , Rema Raman ^(1,2) , Ronald Thomas ⁽²⁾ , Paul Aisen ⁽²⁾ <i>⁽¹⁾ Division of Biostatistics & Bioinformatics, University of California, San Diego, USA, ⁽²⁾ Department of Neurosciences, University of California, San Diego, USA</i>
1.30 - 1.45 p.m	OC16 -	Souvenaid preserves brain function in patients with mild Alzheimer's disease: results from a randomised controlled study <u>Elisabeth CW van Straaten</u> ⁽¹⁾ , Hanneke de Waal ⁽²⁾ , Marieke M.Lansbergen ⁽³⁾ , Philip Scheltens ⁽²⁾ , Cornelis J.Stam ⁽¹⁾ <i>⁽¹⁾ Department of Clinical Neurophysiology, VU University Medical Centre, Amsterdam, The Netherlands, ⁽²⁾ Department of Neurology and Alzheimer Centre, VU University Medical Centre, Amsterdam, The Netherlands, ⁽³⁾ Nutricia Advanced Medical Nutrition, Danone Research, Centre for Specialised Nutrition, Wageningen, The Netherlands</i>
1.45 - 2.00 p.m	OC17 -	The Kinase PKR: a diagnostic and therapeutic target in Alzheimer's disease <u>Jacques Hugon</u> , François Mouton-Liger, Julien Dumurgier, Claire Paquet <i>Memory Center Lariboisiere Hospital APHP, University Diderot Paris France and Institut du Fer a Moulin, Inserm U 839 Paris France</i>
2.00 - 2.15 p.m	OC18 -	Towards Standardization of CSF Biomarkers : A Multi-Site Study Using Validated Assays for A β 42 and Tau <u>Robert M.Umek</u> , David H.Stewart, Jill M.Dunty, Nyssa L.Puskar, Pankaj Oberoi, Jacob N.Wohlstadter <i>MESO SCALE DISCOVERY, Gaithersburg, MD 20877 USA</i>
2.15 - 2.30 p.m	OC19 -	Are neuropsychological tests such as those used in ADNI suitable for long-term trials of cognition enhancers for preclinical Alzheimer <u>Keith Wesnes</u> ^(1,2) , Lon Schneider ⁽³⁾ , Keck School of Medicine of the University of Southern California, USA <i>⁽¹⁾ Bracket Global, Goring, UK, ⁽²⁾ Swinburne University, Melbourne, Australia, ⁽³⁾ Keck School of Medicine of the University of Southern California, USA</i>
2.30 - 2.45 p.m	OC20 -	Phase II trial of Metformin in Amnesic MCI <u>Jose A. Luchsinger</u> ⁽¹⁾ , Jennifer Manly ⁽²⁾ , Jason Steffener ⁽²⁾ , Emilia Bagiella ⁽³⁾ <i>⁽¹⁾ Department of Medicine, Columbia University Medical Center, New York, New York, USA, ⁽²⁾ Gertrude H. Sergievsky Center, Columbia University Medical Center, New York, New York, USA, ⁽³⁾ Biostatistics Center for Clinical Trials Management, Mt. Sinai School of Medicine, New York, NY, USA</i>
2.45 - 3.00 p.m	OC21 -	γ -secretase-linked clinical trials failures: An amyloid cascade hypothesis rebuttal or just experimental pitfalls unmasked? <u>Frédéric Checler</u> , CNRS, Sophia Antipolis, France

3.00 - 4.00 p.m	Symposium 6 : Modeling the Course of AD: Contributions to Better Clinical Trials Chairman : <i>René Spiegel</i> , <i>Memory Clinic, Department of Geriatrics, University Hospital, Basel, Switzerland</i>
3.00 - 3.20 p.m	<ul style="list-style-type: none"> • Predicting Progression of AD : Understanding the Variance Rachel S.Doody ⁽¹⁾, <i>Chan Wen</i> ⁽²⁾, <i>Pavlik Valory</i> ⁽¹⁾, <i>Paul Massman</i> ^(1,3), <i>Eveleen Darby</i> ⁽¹⁾, <i>Susan Rountree</i> ⁽¹⁾ ⁽¹⁾ Baylor College of Medicine - Department of Neurology, Houston, Texas, USA, ⁽²⁾ University of Texas Health Science Center, Houston, Texas, USA, ⁽³⁾ University of Houston, Texas, USA
3.20 - 3.40 p.m	<ul style="list-style-type: none"> • Creating a New Composite Score for Optimizing Responsiveness to Decline in Early AD and Very Early AD <i>Suzanne Hendrix</i>, <i>Pentara Corporation, Salt Lake City, UT, USA</i>
3.40 - 4.00 p.m	<ul style="list-style-type: none"> • The Placebo Group Simulation Approach (PGSA): An Alternative to Long-term Placebo-controlled Trials René Spiegel ⁽¹⁾, <i>Manfred Berres</i> ⁽²⁾, <i>André R.Miserez</i> ⁽³⁾, <i>Andreas U.Monsch</i> ⁽¹⁾ ⁽¹⁾ Memory Clinic, Department of Geriatrics, University Hospital, Basel, Switzerland, ⁽²⁾ University of Applied Sciences Koblenz, RheinAhr Campus, Remagen, Germany, ⁽³⁾ Diogene Laboratories Inc., Reinach, Switzerland
4.00 - 4.30 p.m	<i>Coffee Break and poster sessions</i>
4.30 - 5.30 p.m	Symposium 7 : MAPT (Multidomain Alzheimer Preventive Trial) Imaging (MRI, FDG-PET, amyloid-PET) data Chairmen : <i>Bruno Vellas</i> ⁽¹⁾ , <i>Jacques Touchon</i> ⁽²⁾ , <i>Mike Weiner</i> ⁽³⁾ ⁽¹⁾ Toulouse University Hospital, France, ⁽²⁾ Montpellier University Hospital, Inserm U1061, Montpellier, France, ⁽³⁾ University of California San Francisco, USA
4.30 - 4.45 p.m	<ul style="list-style-type: none"> • MAPT Trial Design <i>Bruno Vellas</i>, <i>Toulouse University Hospital, France, Gerontopole, inserm U 1027</i>
4.45 - 5.00 p.m	<ul style="list-style-type: none"> • One year longitudinal study of FDG-PET in MAPT <i>Thierry Voisin</i>, <i>Toulouse University Hospital, France</i>
5.00 - 5.15 p.m	<ul style="list-style-type: none"> • High prevalence of amyloid positive PET in pre frail and frail older adults <i>Pierre Payoux</i>, <i>Toulouse University Hospital, France</i>
5.15 - 5.30 p.m	<ul style="list-style-type: none"> • M.R.I. Data at baseline <i>Carole Dufouil</i>, <i>Bordeaux University Hospital, France</i>

Tuesday, October 30th *Continued*

5.30 - 6.00 p.m		KEYNOTE 2 New design for symptomatic treatment of AD <i>Lon S. Schneider, University of Southern California, Los Angeles, CA, USA</i>
6.00 p.m		ORAL COMMUNICATIONS Moderators : <i>S.Andrieu, R.Spiegel</i>
6.00 - 6.15 p.m	OC22 - Statistical Power of a Preventive Trial in Alzheimer's Disease, about the GuidAge Study <u>Bruno Scherrer</u> ⁽¹⁾ , <u>Philippe Garnier</u> ⁽²⁾ , <u>Hélène Mathieux-Fortunet</u> ⁽²⁾ , <u>Sandrine Andrieu</u> ⁽³⁾ , <u>Bruno Vellas</u> ⁽⁴⁾ ⁽²⁾ <i>Ipsen, Boulogne, France</i> ⁽³⁾ <i>Inserm UMR 1027, Toulouse, France</i> ⁽⁴⁾ <i>Gérontopole, Toulouse, France</i>	
6.15 - 6.30 p.m	OC23 - Detecting And Intervening In Mild Cognitive Impairment: Is It Cost Effective ? <u>Jennifer H Barnett</u> ^(1,2) , <u>Lily Lewis</u> ⁽³⁾ , <u>Andrew Blackwell</u> ^(1,2) , <u>Matthew Taylor</u> ⁽³⁾ ⁽¹⁾ <i>Cambridge Cognition, Cambridge UK</i> ⁽²⁾ <i>University of Cambridge Department of Psychiatry, UK</i> ⁽³⁾ <i>York Health Economics Consortium, University of York, UK</i>	

Wednesday, October 31st

8.00 - 9.00 a.m		UPDATE ON CLINICAL TRIALS 3 Moderators : <i>P.Robert, M.Weiner</i>
8.00 - 8.20 a.m	8 - Challenges and Perspectives in Designing and Conducting Studies in Alzheimer's disease (AD) <u>Christian Yavorsky</u> , <i>CROnos CCS, Hamilton, NJ, USA</i>	
8.20 - 8.40 a.m	9 - Improving Detection of Treatment Effect Through Better Methodology <u>William R. Shankle</u> ^(1,2) , <u>Alireza Atri</u> ^(3,4) , <u>Michael Rafij</u> ^(5,6,7) ⁽¹⁾ <i>Shankle Clinic, Newport Beach, CA USA</i> , ⁽²⁾ <i>Memory and Cognitive Disorders Program, Hoag Neurosciences Institute; Medical Care Corporation; Dept of Cognitive Sciences, University of California at Irvine, CA, USA</i> , ⁽³⁾ <i>Clinical Programs, GRECC, ENRM VA Bedford Medical Center, USA</i> , ⁽⁴⁾ <i>Memory Disorders Unit & MA Alzheimer's Disease Research Center; Massachusetts General Hospital; Harvard Medical School, USA</i> , ⁽⁵⁾ <i>Memory Disorders Clinic at University of California at San Diego Perlman Ambulatory Care Center, San Diego, CA, USA</i> , ⁽⁶⁾ <i>Dept of Neurosciences at the University of California, San Diego, Alzheimer's Disease Cooperative Study, USA</i> , ⁽⁷⁾ <i>Shiley-Marcos Alzheimer's Disease Research Center, USA</i>	
8.40 - 9.00 a.m	10 - How imaging is used to select subjects and determine therapeutic and adverse effects in Alzheimer's treatment trials <i>Michael Weiner, University of California San Francisco, USA</i>	

Wednesday, October 31st

9.00 - 10.00 p.m		ORAL COMMUNICATIONS Moderator : <i>C.Reed, F.Palermi</i>
9.00 - 9.15 a.m	OC24 -	Clinical trials and Ethical issues: new challenges, new perspectives <u>Federico Palermi</u> , Catherine Pastor, <i>AMPA, Monte Carlo</i>
9.15 - 9.30 a.m	OC25 -	A Multi-Center Clinical Trial to Validate Event-Related Potentials as Useful Biomarkers for Early Detection of Alzheimer's disease <u>K.C.Fadem</u> ⁽¹⁾ , Marco Cecchi ⁽¹⁾ , Sarah Berg ⁽¹⁾ , Charles Smith ⁽²⁾ , Gregory Jicha ⁽²⁾ , Paul Solomon ⁽³⁾ , Carl Sadowsky ⁽⁴⁾ , P.Murali Doraiswami ⁽⁵⁾ , Steven Arnold ⁽⁶⁾ ⁽¹⁾ <i>Neuronetrix, KY, USA</i> , ⁽²⁾ <i>Neurology Department, University of Kentucky, KY, USA</i> , ⁽³⁾ <i>The Memory Clinic, VT, USA</i> , ⁽⁴⁾ <i>Premiere Research Institute, FL, USA</i> , ⁽⁵⁾ <i>Psychiatry Department, Duke University, NC, USA</i> , ⁽⁶⁾ <i>Psychiatry Department, University of Pennsylvania, PA, USA</i>
9.30 - 9.45 a.m	OC26 -	Positive effects on cognitive and clinical function in mild to moderate alzheimer's disease patients with a selective alpha-7 nicotinic partial agonist: interpretation of effects based on a pk/pd model <u>DC.Hilt</u> ⁽¹⁾ , M.Gawryl ⁽¹⁾ , G.Koenig ⁽¹⁾ , N.Dgetluck ⁽¹⁾ , J.Harrison ⁽²⁾ , H.J.Moebius ⁽¹⁾ , G.Loewen ⁽¹⁾ , EVP-6124-010 Study Group ⁽¹⁾ <i>EnVivo Pharmaceuticals, Watertown, MA, USA</i> , ⁽²⁾ <i>Metis Cognition Ltd, UK</i>
9.45 - 10.00 a.m	OC27 -	Stability of symmetric atrophy measurement algorithms: impacts on interim and final analyses <u>D.M.Cash</u> ^(1,2) , J.Bartlett ^(1,3) , S.Finnegan ⁽¹⁾ , K.K.Leung ⁽¹⁾ , G.R.Ridgway ⁽⁴⁾ , M.J.Cardoso ⁽¹⁾ , J.M.Schott ⁽¹⁾ , S.Ourselin ^(1,2) , N.C.Fox ⁽³⁾ ⁽¹⁾ <i>Dementia Research Centre, UCL Institute of Neurology, University College London The Alzheimer's Disease Neuroimaging Initiative</i> , ⁽²⁾ <i>Centre for Medical Image Computing, Department of Computer Science, University College London</i> , ⁽³⁾ <i>Department of Medical Statistics, London School of Hygiene and Tropical Medicine</i> , ⁽⁴⁾ <i>Wellcome Trust Centre for Neuroimaging, UCL Institute of Neurology</i>
10.00 - 10.30 a.m		<i>Coffee Break and poster sessions</i>
10.30 - 12.30 p.m		ORAL COMMUNICATIONS Moderator : <i>N.Coley, A.Gabelle</i>
10.30 - 10.45 a.m	OC28 -	pS422 Tau-immunotherapy in THY-Tau22 transgenic mice <u>Luc Buée</u> , Laetitia Troquier, Philippe Lassalle, David Blum, <i>Inserm U837 - Alzheimer & Tauopathies, Faculty of Medicine, Research Department, Lille, France</i>
10.45 - 11.00 a.m	OC29 -	Outpatients with severe Alzheimer's disease participating in an observational study: Baseline results of the GERAS study <u>Catherine Reed</u> ⁽¹⁾ , Josep Maria Argimon ⁽²⁾ , Mark Belger ⁽¹⁾ , Giuseppe Bruno ⁽³⁾ , Richard Dodel ⁽⁴⁾ , Michael Happich ⁽⁵⁾ , Josep Maria Haro ⁽⁶⁾ , Roy W.Jones ⁽⁷⁾ , Bruno Vellas ⁽⁸⁾ , Anders Wimo ⁽⁹⁾ ⁽¹⁾ <i>Eli Lilly and Company Limited, Windlesham, UK</i> , ⁽²⁾ <i>Servei Català de la Salut, Barcelona, Spain</i> , ⁽³⁾ <i>University of Rome «Sapienza», Rome, Italy</i> , ⁽⁴⁾ <i>Philipps-University, Marburg, Germany</i> , ⁽⁵⁾ <i>Lilly Deutschland GmbH, Bad Homburg, Germany</i> , ⁽⁶⁾ <i>Parc Santari Sant Joan de Déu, CIBERSAM, Universitat de Barcelona, Barcelona, Spain</i> , ⁽⁷⁾ <i>The RICE Centre, Royal United Hospital, Bath, UK</i> , ⁽⁸⁾ <i>Toulouse University Hospital, Toulouse, France</i> , ⁽⁹⁾ <i>Karolinska Institute, Stockholm, Sweden</i>

Wednesday, October 31st *Continued*

ORAL COMMUNICATIONS - Continued

11.00 - 11.15 a.m	OC30 -	Level of education, risk of Alzheimer's disease, and cognitive decline in the GuidAge prevention trial <u>Nicola Coley</u> ^(1,2) , Grégory Guernec ⁽¹⁾ , Bruno Vellas ^(1,2,3) , Sandrine Andrieu ^(1,2,3,4) and the Guidage study group ⁽¹⁾ INSERM, U1027, Toulouse, France, ⁽²⁾ Université de Toulouse III, Toulouse, France, ⁽³⁾ Gérontopôle, Centre Hospitalier Universitaire de Toulouse, Toulouse, France, ⁽⁴⁾ Département d'Epidémiologie et Santé Publique, Centre Hospitalier Universitaire de Toulouse, Toulouse, France
11.15 - 11.30 a.m	OC31 -	Repurposing cardiovascular drugs as Alzheimer's disease modifying agents <u>Giulio Maria Pasinetti</u> ⁽¹⁾ , Paul Rosenberg ⁽²⁾ ⁽¹⁾ Department of Neurology, Mount Sinai School of Medicine, New York, USA, ⁽²⁾ Department of Psychiatry, Johns Hopkins School of Medicine, Baltimore, USA
11.30 - 11.45 a.m	OC32 -	Comparison of interim results after one and two years of treatment of ASCOMALVA TRIAL between a cholinesterase inhibitor and choline alphoscerate on cognitive deficits in Alzheimer's disease associated with cerebrovascular impairment <u>Francesco Amenta</u> ⁽¹⁾ , Anna Carotenuto ⁽¹⁾ , Raffaele Rea ⁽¹⁾ , Enea Traini ⁽¹⁾ , Angiola M.Fasanaro ⁽²⁾ ⁽¹⁾ Center of Clinical Research, telemedicine and telepharmacy, University of Camerino, Italy, ⁽²⁾ Evaluation Department for Alzheimer's Disease and other Neurodegenerative disease, National University Hospital A. Cardarelli, Napoli, Italy
11.45 - 12.00 p.m	OC33 -	Translating Cognitive Performance to Functional Ability in Alzheimer's, Lewy Body, Cerebrovascular, and Frontal Lobe Diseases <u>William R.Shankle</u> ^(1,3,5) , Junko Hara ^(1,2) , Barry Reisberg ⁽⁴⁾ , Michael Lee ⁽⁵⁾ , James Pooley ⁽⁵⁾ ⁽¹⁾ Shankle Clinic, Newport Beach, CA, USA, ⁽²⁾ Medical Care Corporation, Newport Beach, CA, USA, ⁽³⁾ Hoag Neurosciences Institute, Newport Beach, CA, USA, ⁽⁴⁾ Aging & Dementia Research Center, New York University, New York, USA, ⁽⁵⁾ Dept. of Cognitive Sciences, University of California at Irvine, Irvine, CA, USA
12.00 - 12.15 p.m	OC34 -	Deregulation of Cdk5 activity and neurodegeneration induced by hyperphosphorylation of neuronal cytoskeletal proteins is inhibited by peptides derived from its activator p35; A novel approach to rescue the pathology induced by deregulation of kinases <u>Harish C.Pant</u> , NINDS / NIH, Bethesda, MD, USA
12.15 - 12.30 p.m	OC35 -	Disease-Modifying Power Estimation for TauRx Global Phase 3 Trial Program in AD with Tau-Aggregation Inhibitor LMTX <u>Claude M.Wischik</u> , Damon J.Wischik, TauRx Therapeutics, USA

SCIENTIFIC POSTERS

For your convenience posters are classified by themes and numbered accordingly :

THEMES		
Clinical Trials : Methodological aspects	P1 through P12	p.16 to 17
Clinical trials Assessment Tools	P13 through P23	p.17 to 18
Therapeutic trials in AD	P24 through P35	p.18 to 19
Fundamental Research for AD clinical trials	P36 through P42	p.20
Biomarkers/Imaging in AD clinical trials	P43 through P63	p.21 to 23
Nutrition and Alzheimer's Disease	P64 through P71	p.23 to 24
Miscellaneous in clinical trials	P72 through P88	p.24 to 25

SCIENTIFIC POSTERS

Clinical Trials: Methodological aspects

- P1** Meta-Analysis for Behavioral and Neuropsychiatric Symptoms in Alzheimer's disease (AD) Clinical Trials
Kaori Ito, Sima Ahadiieh, Peter Lockwood, Thomas Tensfeldt, Brian Corrigan
Pfizer Inc, Primary Care Business Unit, Groton, CT, USA
- P2** Characteristics of Alzheimer's patients participating in research within the French National Alzheimer Bank Network
Karim Tifratene (1,2), Franck Leduff (1), Christian Pradier (1), Malthilde Laye (2), Roland Chevrier (1), Philippe Robert (2)
(1) Public Health Department, Nice University Hospital, France, (2) CMRR, EA CobTeK, Nice University
- P3** Clinical background of participants of clinical trials on Alzheimer's disease is different from that of actual patients.
Motoki Yutani (1), Megumi Takahashi (2), Satoru Oishi (2), Kumiko Shichijou (3), Hitoshi Miyaoka (2)
(1) Kitasato University Graduate School of Medical Sciences, (2) Department of Psychiatry, Kitasato University School of Medicine, (3) Clinical trial center, Kitasato University East Hospital
- P4** Predictors of progression from mild cognitive impairment to Alzheimer's disease in the placebo arm of a clinical trial population
Niels D.Prins (1), Wiesje M.Van der Flier (1), H.Robert Brashear (2), Frederik Barkhof (1,3), Philip Scheltens (1)
*(1) Alzheimer Center and department of Neurology, VU University Medical Centre, Amsterdam, the Netherlands, (2) Janssen Alzheimer Immunotherapy Research and Development, South San Francisco, CA, USA
 (3) Alzheimer Centre, Department of Radiology, and Image Analysis Centre, VU University Medical Centre, Amsterdam, the Netherlands*
- P5** The SKT short cognitive performance test in patients with dementia: Cross-sectional and longitudinal correlations with activities of daily living in two independent clinical trials
R.Hoerr (1), H.Lehfeld (2), S.Schlaefke (1)
(1) Clinical Research Department, Dr. Willmar Schwabe GmbH & Co. KG, Karlsruhe, Germany, (2) Department of Psychiatry and Psychotherapy, Nuremberg Hospital, Nuremberg, Germany
- P6** Innovative Digital Patient Recruitment Strategies in Prodromal AD trials
Lynne Hughes (1), Amir Kalali (1), Cathy Vanbelle (1), Elisa Cascade (1)
Quintiles, Inc
- P7** Cognitive impairment, dementia and «good performers»: prevalence in the population study «Invece.Ab»
Antonio Guaita (1), Simona Abbondanza (1), Mauro Colombo (2), Annalisa Davin (1), Gianluigi Forloni (3), Letizia Polito (1), Roberta Vaccaro (1), Eleonora Valle (1), Silvia Vitali (2), Virginia V Ferretti (4), Simona Villani (4)
(1) GolgiCenci Foundation, Abbiategrasso (Milan), Italy, (2) Geriatric Institute "C. Golgi", Abbiategrasso (Milan), Italy, (3) Department of Neuroscience, Mario Negri Institute for Pharmacological Research, Milan, Italy, (4) Department of Health Sciences, Section of Medical Statistics and Epidemiology, University of Pavia, Pavia, Italy
- P8** The Placebo Group Simulation Approach (PGSA): Its Validity when Applied to Data from Alzheimer Patients in the ADNI 1 Database
René Spiegel (1), Manfred Berres (2), Andreas U.Monsch (1)
(1) University Hospital, Department of Geriatrics, Memory Clinic, Basel Switzerland, (2) University of Applied Sciences Koblenz, RheinAhr Campus, Remagen Germany
- P9** Intra-individual variability (IIV) vs. total scores as the target outcome in clinical trials
Rochelle E.Tractenberg (1,2), Futoshi Yumoto (2,3), Paul S.Aisen (4)
(1) Departments of Neurology, Biostatistics, Bioinformatics & Biomathematics, and Psychiatry, Georgetown University Medical Center, Washington, D.C., USA, (2) Collaborative for Research on Outcomes and Metrics, USA, (3) University of Maryland University College, College Park, MD, USA, (4) Department of Neurology, University of California, San Diego, La Jolla, CA, USA
- P10** Review and Analysis of Rater Errors on the Neuropsychological Test Battery (NTB) in an International Alzheimer's Disease Clinical Trial
Lisle Kingery (1), Jenn Cromer (1,2), Brian Merry (1), Alison Miner (1), Judy Jaeger (1), Amy Veroff (1), Paul Maruff (3)
(1) Cogstate, New Haven, CT, USA, (2) Hospital for Special Care, New Britain, CT, USA, (3) CogState, Melbourne, Victoria, Australia

Clinical Trials: Methodological aspects - *Continued*

- P11** INROADS: International Registry of Alzheimer's Disease Patients
RA.Stern (1), PN.Tariot (2), HM.Arrighi (3), LA.Lacey (4), GA.Rippon (5), V.Menon (6), MT.Ropacki (3)
 (1) Boston University, Boston, MA, (2) Banner Alzheimer's Institute, Phoenix, AZ, (3) Janssen Alzheimer Immunotherapy Rsearch & Development, LLC, South San Francisco, CA, USA, (4) Janssen Alzheimer Immunotherapy, Dublin, Ireland, (5) Pfizer Inc, Collegeville, PA, (6) Outcome Sciences, Cambridge, MA
- P12** Neuropsychiatric Syndromes in Mild versus Moderate AD: Factor Analysis from a Clinical Trial Population
Constantine Lyketsos (1), Susan Abushakra (2), Gerald Crans (2), Chito Hernandez (2), Sandrine Andrieu (3), Bruno Vellas (3)
 (1) Johns Hopkins University, Baltimore, MD, USA, (2) Elan Pharmaceuticals, Inc., South San Francisco, CA, USA, (3) University of Toulouse, Toulouse, France

Clinical trials Assessment Tools

- P13** Patient and carer views on clinical trials using immunotherapy for prodromal Alzheimer's disease
Vanessa Lawrence (1), James Pickett (2), Joanna Murray (1)
 (1) Institute of Psychiatry, King's College London, UK, (2) Alzheimer's Society, London, UK
- P14** Can Screen to Baseline MMSE Variability in AD Clinical Trials Affect Primary Outcome Measurement?
David S.Miller (1), Yanhui Xu (1), Priscilla Samuelson (1), David Henley (2), Gopalan Sethuraman (2), Karen Sundell (2)
 (1) Brackett, Wayne, PA, USA, (2) Eli Lilly & Co., Indianapolis, IN, USA
- P15** ADAS-Cog is an effective tool for Cognitive Research in Southeast Asia
Nagaendran Kandiah, Ivane Chew, Shan Huang, Amanda Ng
 Department of Neurology, National Neuroscience Institute, Singapore
- P16** National Institutional Review Board for Neurodegenerative Diseases (NIRB-ND) for the United States
Ara S.Khachaturian (1), Peter J.Snyder (2), Maria Carrillo (3), David S.Knopman (4)
 (1) The Campaign to Prevent Alzheimer's Disease by 2020 (PAD2020), Rockville, Maryland, USA, (2) Lifespan Hospitals & Alpert Medical School of Brown University, Providence, Rhode Island, USA, (3) The Alzheimer's Association, Chicago Illinois, USA, (4) The Mayo Clinic & Alzheimer's Disease Cooperative Study, Rochester, Minnesota, USA
- P17** Assessment of consent ability in Alzheimer's disease during research: a pilot study of a French questionnaire
Nathalie Philippi (1), Chloé Chaulet-Dalard (2), Isabelle Migeon-Duballet (3), Barbara Jung (1), Olga Uspenskaya (4), H  l  ne Mollion (5), Karim Bennys (2), Lynda Touati (2), Audrey Gabelle-Deloustal (2), Jacques Touchon (2,7), Benjamin Cretin (1), Marc Paccalin (3), Catherine Martin-Hunyadi (1), Fr  d  ric Blanc (1), Jean-Christophe Weber (1)
 (1) Strasbourg University Hospital, France, (2) Montpellier University Hospital, France, (3) Poitiers University Hospital, France, (4) IM2A, Piti  -Salp  tri  re University Hospital, Paris, France, (5) Lyon University Hospital, France, (6) IRIST – EA 3424, Strasbourg University, France, (7) INSERM U1061, Montpellier, France
- P18** The Memory and Attention Test (MAT): a computer-based test for studies in dementia
Georg Adler (1), Miriam Bektas (1), Nadja Baumgart (2), Yvonne Lembach (1)
 (1) Institut fuer Studien zur Psychischen Gesundheit (ISPG), Mannheim, Germany, (2) Dynamikos GmbH, Mannheim, Germany
- P19** Establishing threshold scores and profiles of cognitive impairment for the Alzheimer's Disease Assessment Scale (ADAS-Cog) for patients with probable Mild Cognitive Impairment (MCI)
C.Yavorsky (1,2), G.DiClemente (1), M.Opler (2), A.Khan (2,3), S.Jovic (2), B.Rothman (2)
 (1) CROnos CCS, Hamilton, NJ, (2) ProPhase LLC, New York, NY, (3) Nathan S.Kline Institute for Psychiatric Research

SCIENTIFIC POSTERS

Clinical trials Assessment Tools - *Continued*

- P20** Testing the dynamic model of Alzheimer's disease in two European memory clinics
Giovanni B. Frisoni (1), Anna Paola Prestia (1), Anna Caroli (1,2), Wiesje M. Van der Flier (3,4), Rik Ossenkoppele (3,5), Bart N.M. Van Berckel (5), Frederik Barkhof (6), Charlotte E. Teunissen (7), Philip Scheitens (3), Translational Outpatient Memory Clinic – TOMC - Working Group (1) LENITEM – Laboratory of Epidemiology Neuroimaging and Telemedicine, IRCCS Centro San Giovanni di Dio FBF, Brescia, Italy, (2) Medical Imaging Unit, Biomedical Engineering Department, Mario Negri Institute for pharmacological research, Bergamo, Italy, (3) Alzheimer Center and dept of Neurology, VU University Medical Center, Amsterdam, The Netherlands, (4) Dept of Epidemiology & Biostatistics, VU University Medical Center, Amsterdam, The Netherlands, (5) Dept of Nuclear Medicine and PET research, VU University Medical Center, Amsterdam, The Netherlands, (6) Dept of Radiology, VU University Medical Center, Amsterdam, The Netherlands, (7) Dept of Clinical Chemistry, VU University Medical Center, Amsterdam, The Netherlands
- P21** ADAS-EXEC: A cognitive composite derived from the ADAS-cog and NTBA
John Harrison (1,2), Jill Altman (3), Geoffrey Kempler (4), Dianne Angus (4), Caroline Herd (4), Steve Targum (5), Jeffrey Cummings (6), Craig Ritchie (2) (1) Metis Cognition Ltd., Kilmington Common, UK, (2) Freelance Statistician, Redhill, UK, (4) Prana Biotechnology Ltd., Melbourne, Australia, (5) ClintaraInc, Boston, US, (6) Cleveland Clinic's Lou RuvoCenter for Brain Health, Las Vegas, Nevada
- P22** Benefits of an Item Response Theory Based Analysis of ADAS-cog Assessments
Sebastian Ueckert (1), Elodie L.Plan (2), Kaori Ito (3), Mats O.Karlsson (1), Brian Corrigan (3), Andrew C.Hooker (1) (1) Department of Pharmaceutical Biosciences, Uppsala University, Uppsala, Sweden, (2) Metrum Research Group, Tariffville, CT, USA, (3) Pfizer Inc, Primary Care Business Unit, Groton, CT, USA
- P23** Pharmacometric Modeling of Clinical ADAS-cog Assessment Data using Item Response Theory
Sebastian Ueckert (1), Elodie L.Plan (2), Kaori Ito (3), Mats O.Karlsson (1), Brian Corrigan (3), Andrew C.Hooker (1) (1) Department of Pharmaceutical Biosciences, Uppsala University, Uppsala, Sweden, (2) Metrum Research Group, Tariffville, CT, USA, (3) Pfizer Inc, Primary Care Business Unit, Groton, CT, USA

Therapeutic trials in AD

- P24** Long-term administration of active immunotherapy CAD106 in Phase IIa open-label extension studies in Alzheimer Patients
A. Graf (1), ME.Riviere (1), M.Farlow (2), N.Andreasen (3), P.Quarg (1), A.Caputo (1), I.Vostiar (1), B.Winblad (3), JM.Orgogozo (4) (1) Novartis Pharma AG, Basel, Switzerland, (2) Indiana University, Indianapolis, USA, (3) Karolinska Universitetssjukhuset, Huddinge, Sweden, (4) Université de Bordeaux, France
- P25** Evaluating the Cognitive Effects of Donepezil 23 mg/d in Moderate and Severe Alzheimer's Disease: A Patient Subgroup Analysis
Marwan Sabbagh (1), Jeffrey Cummings (2), Daniel Christensen (3), Rachelle Doody (4), Martin Farlow (5), Joan Mackell (6), Randi Fain (7) (1) Banner Sun Health Research Institute, Sun City, AZ, USA, (2) Cleveland Clinic Lou Ruvo Center for Brain Health, Las Vegas, NV, USA, (3) University of Utah, Salt Lake City, UT, USA, (4) Baylor College of Medicine, Houston, TX, USA, (5) Indiana University School of Medicine, Indianapolis, IN, USA, (6) Pfizer Inc, New York, NY, USA, (7) Eisai Inc., Woodcliff Lake, NJ, USA
- P26** Donepezil 23 mg/d for Moderate to Severe Alzheimer's Disease: Assessing Subdomains of the Severe Impairment Battery
Steven Ferris (1), Jeffrey Cummings (2), Daniel Christensen (3), Rachelle Doody (4), Martin Farlow (5), Marwan Sabbagh (6), Joan Mackell (7), Randi Fain (8) (1) Alzheimer Disease Center, New York University Langone Medical Center, New York, NY, USA, (2) Cleveland Clinic Lou Ruvo Center for Brain Health, Las Vegas, NV, USA, (3) University of Utah, Salt Lake City, UT, USA, (4) Baylor College of Medicine, Houston, TX, USA, (5) Indiana University School of Medicine, Indianapolis, IN, USA, (6) Banner Sun Health Research Institute, Sun City, AZ, USA, (7) Pfizer Inc, New York, NY, USA, (8) Eisai Inc., Woodcliff Lake, NJ, USA
- P27** Simvastatin versus atorvastatin: which is the optimal choice to prevent Alzheimer's disease?
*Saleta Sierra, Javier S.Burgos
 BioPharma Division, Neuron Bio, Parque Tecnológico de Ciencias de la Salud, Granada, Spain*

Therapeutic trials in AD - *Continued*

- P28** A Nicotinic-alpha-7 Partial Agonist as adjunctive therapy to stable Donepezil
Donna Masterman (1), Tari Awipi (1), Elizabeth Ashford (1), Stephane Nave (1), Kisoook Yoo (1), Bruno Vellas (2), Luca Santarelli (1)
 (1) CNS DTA, F. Hoffmann-La Roche, Ltd., Basel, Switzerland, (2) Clinic of Internal Medicine and Gerontology, University Hospital Center, Toulouse, France
- P29** Effects of ELND005 (Scyllo-inositol) on Neuropsychiatric Symptoms (NPS) in Mild / Moderate AD: Correlations of ELND005 Exposures to Neuropsychiatric Outcomes in a 78-Week Phase 2 Study
Earvin Liang (1), Jonathan Wagg (2), Matthias Kurth (1), Susan Abushakra (1)
 (1) Eian Pharmaceuticals, Inc., South San Francisco, CA, USA, (2) Pharsight/Cartara Corporation, St. Louis, MO, USA
- P30** The Study of Pharmacological Efficacy of Dry Mulberry Burirum-60 in Alzheimer's Disease
Buavaroon Srichaiku, Waraporn Sutthisa Hospital, Thailand
- P31** Cerebrolysin, a novel drug for the treatment of Alzheimer's Disease. An Experimental Study using nanowired delivery
Hari S Sharma (1), Rudy J Castellani (2), Mark A Smith (3), Z Ryan Tian (4), Dafin F Muresanu (5), Herbert Mössler (6), Aruna Sharma (1)
 (1) Cerebrovascular Research Laboratory, Department of Surgical Sciences, Anesthesiology & Intensive Care Medicine, University Hospital, Uppsala University, Sweden, (2) Department of Pathology, University of Maryland, Baltimore, MD, USA, (3) Department of Pathology, Case Western Reserve University, Cleveland, Ohio USA, (4) Department of Chemistry & Biochemistry, University of Arkansas Fayetteville, Fayetteville, AR, USA, (5) Department of Neurosciences, University of Medicine & Pharmacy, University Hospital, Cluj-Napoca, Romania, (6) Ever Neuro Pharma, Oberburgau, Austria
- P32** Implication of Integrative Treatment for Real-Life Geriatric Patients with Depression, Dementia and Multiple Chronic Diseases: A 60-Month Follow-Up of a Naturalistic Study
Gjumrakch Aliev (1,2), Valentin Bragin (3), Ramon Cacabelos (4)
 (1) Department of Health Science and Healthcare Administration, University of Atlanta, Atlanta, GA USA, (2) GALLY™ International Biomedical Research Consulting LLC, San Antonio, TX, USA, (3) Stress Relief and Memory Training Center, Brooklyn, New York, NY USA, (4) EuroEspes Biomedical Research Center, Institute for CNS Disorders and Genomic Medicine & Camilo José Cela University, La Coruña, Spain
- P33** A pharmacogenetics-supported clinical trial to delay onset of mild cognitive impairment (MCI) due to Alzheimer's disease (AD)
Allen D. Roses (1,2), Kathleen A. Welsh-Bohmer (2), Daniel K. Burns (1), Carl Chiang (1), Donna G. Crenshaw (1,2), Michael W. Lutz (1,2), Craig A. Metz (1), Ann M. Saunders (1,2), Stephen Brannan (3), Manoj Malhotra (3)
 (1) Zinfandel Pharmaceuticals, Inc., Durham, NC, USA, (2) Duke Bryan ADRC, Durham, NC, USA, (3) Takeda Global Research & Development Center, Inc., Deerfield, IL, USA
- P34** Cognitive stimulation and APOE genotype in non-demented elderly subjects: a randomized controlled study (RCT)
Gianluigi Forloni (1), Letizia Polito (2), Annalisa Davin (2), Simona Abbondanza (2), Roberta Vaccaro (2), Eleonora Valle (2), Antonio Guaita (2), Mauro Colombo (3), Silvia Vitali (3), Virginia Valeria Ferretti (4), Simona Villani (4)
 (1) Department of Neuroscience, Mario Negri Institute for Pharmacological Research, Milan, Italy, (2) GolgiCenci Foundation, Abbiategrasso (Milan), Italy, (3) Geriatric Institute "C. Golgi", Abbiategrasso (Milan), Italy, (4) Dep. of Health Sciences, Biostatistics and Clinical Epidemiology unit, Pavia University, Pavia, Italy
- P35** TauRx Global Phase 3 Clinical Trial of Tau-Aggregation Inhibitor Therapy with LMTX (TM)
Claude M. Wischik
 University of Aberdeen, UK, TauRx Therapeutics

SCIENTIFIC POSTERS

Fundamental Research for AD clinical trials

- P36** Dopamine D2-agonist Rotigotine effects on cortical excitability and central cholinergic transmission in Alzheimer's disease patients.
Alessandro Martorana (1), Giacomo Koch (2)
(1) Clinica Neurologica- Dipartimento di Neuroscienze, Centro di Riferimento per la Malattia di Alzheimer- Università di Roma, Italy, (2) Fondazione Santa Lucia IRCCS, Rome, Italy
- P37** Cellular and Animal Models for High Through-Put Screening of Therapeutic Agents for the Treatment of Diseases of Aging in General and Alzheimer's Disease in Particular
Jordan L. Holtzman
Departments of Pharmacology and Medicine and Division of Environmental Health Sciences, University of Minnesota, Minneapolis, USA
- P38** Potential of Human analogue of Morris Water Maze in translational medicine and the assessment of therapeutic response
John Harrison (2,4,5), Jan Laczó (1,2), Manfred Windisch (2,3), Jakub Hort (1,2)
(1) Memory Disorders Clinic, Department of Neurology, Charles University in Prague, 2nd Medical Faculty and University Hospital Motol, Prague, Czech Republic, (2) Polyhymnia Translational Research, London, UK, (3) JSW-Lifesciences GmbH, Grambach-Graz, Austria, (4) Metis Cognition Ltd., Kilmington, UK, (5) Dept. of Medicine, Imperial College, London, UK
- P39** Voluntary exercise suppresses enhanced tau pathology by high calorie diet in a tauopathy mouse model
Yasumasa Yoshiyama
Department of Neurology and Clinical Research Center, Chiba East National Hospital, Chiba 260-8712, Japan
- P40** Monoclonal antibody generation based on the humanized yeast model system: improving tau diagnostics
Jeff Van den Brande (1,2), Joëlle Rosseels (1), Peter Borghaeef (3), Dirk Jacobs (2), Marleen Michels (1), Luc Buée (4), Fred Van Leuven (3), Eugeen Vanmechelen (2), Joris Winderickx (1)
(1) Functional Biology, KU Leuven, Heverlee, Belgium, (2) ADx Neurosciences, Ghent, Belgium, (3) LEGTEGG, KU Leuven, Belgium, (4) Inserm, U837, Alzheimer & Tauopathies, Lille, France ; Université Lille Nord de France, JP Aubert Research Centre, IMPRT, Lille, France ; UDSL, Faculté de Médecine-Pole Recherche, Lille, France ; CHU-Lille, Lille, France
- P41** Examining start up for a multicenter clinical trial of neurosurgical administration of nerve growth factor gene transfer
Sarah Walter (1), Elizabeth Shaffer-Bacareza (1), Ronelyn Chavez (1), Kristin Woods (1), Genevieve Matthews (1), Paul Aisen (1), Joshua Grill (2)
(1) Alzheimer's Disease Cooperative Study, University of California San Diego, La Jolla, CA, USA, (2) Mary S. Easton Center for Alzheimer's Disease Research, Department of Neurology, David Geffen School of Medicine, University of California, Los Angeles, CA, USA
- P42** Molecular dynamics simulations applying for designing compounds that can be used for treatment of Alzheimer's disease
Maricarmen Hernández (1), Martha Rosales (2), Jose Correa Basurto (1)
(1) Laboratory of Molecular Modeling and Bioinformatics. Escuela Superior de Medicina. IPN, (2) Laboratory of Biophysics and Biocatalysis. Escuela Superior de Medicina. IPN

Biomarkers/Imaging in AD clinical trials

- P43** Changes of Biological Markers and Brain PET Imaging and Clinical Effects of Memantine for Patients with Moderate to Severe Alzheimer's Disease: a 24 Week Double-blind, Randomized, Placebo-Controlled Study
Shifu Xiao (1,2), Tao Wang (1,2), Qiu Huang (3), Kewei Chen (3,4), Eric Reiman (4)
 (1) Geriatric Psychiatry Department, Shanghai Mental Health Center, Shanghai JiaoTong University School of Medicine, Shanghai, China, (2) Alzheimer's Disease and Related Disorder Center, Shanghai JiaoTong University, Shanghai, China, (3) Med-X Institute, Shanghai Jiaotong University, Shanghai, China, (4) Banner Alzheimer's Institute and Banner Good Samaritan Medical Center, PET Center, Phoenix, AZ, USA
- P44** Organotypic hippocampal slices coupled with multi-electrode array: an innovative method to investigate neuroactive compounds in neurodegenerative disorders
Venceslas Duveau, Céline Bouyssieres, Mélanie Langlois, Corinne Roucard
 SynapCell SAS, Biopolis, 5 avenue du Grand Sablon, La Tronche, France
- P45** Development and Application of a Mathematical Model of Modulation of CSF A β 40 in Cerebrospinal Fluid after Treatment with γ and β -secretase inhibitors
Huub Jan Kleijn, Lei Ma, Marissa Dockendorf, Rik de Greef, John Palcza, Jim Kost, Mark Forman, Michael Egan, Julie Stone
 Merck Research Laboratories, Whitehouse Station, NJ
- P46** Definition of Harmonized Protocol for Hippocampal Segmentation
Marina Boccardi (1), Martina Bocchetta (1,2), Liana Apostolova (3), Josephine Barnes (4), George Bartzokis (5), Gabriele Corbetta (1), Charles DeCarli (6), Leyla deToledo-Morrell (7), Michael Firbank (8), Rossana Ganzola (1), Lotte Gerritsen (9), Wouter Henneman (10), Ronald J.Killiany (11), Nikolai Malykhin (12), Patrizio Pasqualetti (2), Jens C.Pruessner (13), Alberto Redolfi (1), Nicolas Robitaille (14), Hilikka Soininen (15), Daniele Tolomeo (1), Lei Wang (16), Craig Watson (17), Henrike Wolf (18), Simon Duchesne (14), Clifford R.Jack Jr (19), Giovanni B.Frisoni (1)
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- P47** Robustness of Automated Hippocampal Volumetry across MR Field strengths and Same-Day Repeat Scans
Robin Wolz (1,2), Adam J Schwarz (3), Peng Yu (3), Pat Cole (4), Daniel Rueckert (1,2), David Raunig (5), Derek Hill (1)
 (1) IXICO Ltd, London, UK, (2) Imperial College London, London, UK, (3) Eli Lilly and Company, Indianapolis IN, USA, (4) Cole Consulting, NJ, USA, (5) Icon Medical Imaging, Warrington PA, USA
- P48** Disentangling the normal aging from the pathological Alzheimer's disease progression on structural MR images
Marco Lorenzi (1,2), Nicholas Ayache (2), Xavier Pennec (2), Giovanni B Frisoni (1), and the Alzheimer's Disease Neuroimaging Initiative (ADNI)
 (1) IRCCS San Giovanni di Dio Fatebenefratelli, Brescia, Italy, (2) Asclepius research project, INRIA Sophia antipolis, France
- P49** Integrating Biomarkers in 12 AD Programs
Amir Kalali, Cathy Vanbelle, Roza Hayduk, Lynne Hughes, Quintiles Inc.
- P50** Contribution of the N400 event related potential in the study of the Primary Progressive Aphasia
Karim Bennys (1), Elise Benattar (1), Gérard Rondouin (2), Sylvie Moritz-Gasser (1), Guillaume Taïeb (3), Jacques Touchon (1,4), Audrey Gabelle-Deloustal (1)
 (1) Department of Neurology, Memory Research Ressource Center for Alzheimer's Disease, University Hospital of Montpellier, France, (2) Neurophysiology Unit, Department of Neurology, University Hospital of Montpellier, France, (3) Department of Neurology, University Hospital of Nîmes, France, (4) INSERM U1061, Montpellier, France

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Biomarkers/Imaging in AD clinical trials - *Continued*

- P51** Supporting evidence of the AD biomarker dynamic model in patients with mild cognitive impairment
Samantha Galluzzi (1), Cristina Geroldi (2), Giovanni Amicucci (3), Luisella Bocchio-Chiavetto (4), Matteo Bonetti (5), Cristian Bonvicini (6), Maria Cotelli (7), Roberta Ghidoni (8), Barbara Paghera (9), Orazio Zanetti (2), Giovanni B Frisoni (1), Translational Outpatient Memory Clinic Working Group
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- P52** Mass Spectrometry Quantitation of Multi-component Protein Biomarker Panels of Alzheimer's Disease
Emma LSchofield (1), Ian Pike (1), Stephan Jung (2), Hans Dieter Zucht (2), Stefan Selzer (2), Christopher Lößner (2), Malcolm A Ward (1)
 (1)Proteome Sciences plc, UK (2)Proteome Sciences R&D GmbH & Co. KG, Frankfurt/Main, Germany
- P53** Investigating Tau and phospho-Tau moieties present within CSF to further biomarker development
Emma L.Schofield (1), Claire L.Russell (1), Christopher Lößner (2), Thorsten Prinz (2), Malcolm A.Ward (1), Ian Pike (1)
 (1) Proteome Sciences plc, UK, (2) Proteome Sciences R&D GmbH & Co. KG, Frankfurt/Main, Germany
- P54** Quantification of cerebral glucose metabolism during normal aging: a comparison of cerebral metabolic rate and spm analysis.
Scott Nugent (1,2), Sébastien Tremblay (1,3,4), Alexandre Castellano (1,2), Mélanie Fortier (1), Eric Turcotte (3,4), Tamas Fulop (1,5), Michèle Allard (6), Stephen Cunnane (1,2,5)
 (1) Research Center on Aging, Sherbrooke, QC, Canada, (2) Department of Physiology and Biophysics, Université de Sherbrooke, QC, Canada, (3) Department of Radiobiology and Nuclear Medicine, Université de Sherbrooke, QC, Canada, (4) Centre for Molecular Imaging, Sherbrooke, QC, Canada, (5) Department of Medicine, Université de Sherbrooke, QC, Canada, (6) Nuclear Medicine Service, Hôpital Pellegrin, UMR-CNRS-5227, EPHE, Bordeaux, France
- P55** MR Imaging Features of ARIA (Amyloid-Related Imaging Abnormalities) in Patients Treated in Phase 2 Clinical Studies of Bapineuzumab
J.Barakos, R.Sperling, S.Salloway, C.Jack, A.Gass, J.Fiebich, D.Tampieri, D.Melançon, Y.Miaux, G.Rippon, R.Black, Y.Lu, HR.Brashear, HM.Arrih, K.Morris, M.Grundman
- P56** Measurement invariance in MRI-derived volumetrics, but not memory tests: implications for longitudinal research of neurodegenerative disease
RE.Tractenberg (1,2), Futoshi Yumoto (2,3)
 (1) Departments of Neurology, Biostatistics, Bioinformatics & Biomathematics, and Psychiatry, Georgetown University Medical Center, Washington, D.C., USA, (2) Collaborative for Research on Outcomes and Metrics, USA, (3) University of Maryland University College, College Park, MD, USA, (4) Department of Neurology, University of California, San Diego, La Jolla, CA, USA
- P57** The distribution of A β 42 among the different blood fractions might help in the early diagnosis of Alzheimer's disease
Pedro Pesini (1,2), Virginia Pérez-Grijalba (1), Mercè Boada (3), Lluís Tàrraga (3), Pablo Martínez-Lage (3), Itziar San-José (2), Manuel Sarasa (1,2)
 (1) Araclon Biotech Ltd., Proteomic Laboratory, CIBIR Logroño, Spain, (2) Araclon Biotech Ltd., I + D Laboratory, Zaragoza, Spain, (3) Alzheimer Research Center and Memory Clinic. Fundació ACE. Institut Català de Neurociències Aplicades. Barcelona, Spain
- P58** Validation of a multi-atlas segmentation technique for the quantification of hippocampal volume - Application as a selection criterion in clinical trials in Alzheimer's Disease
H.J.Yu (1), L.Bracoud (1), J.Schaerer (1), D.Xu (1), F.Roche (1), B.Belaroussi (1), C.Pachai (1), C.DeCarli (2) and the Alzheimer's Disease Neuroimaging Initiative
 (1) BioClinica Inc., Lyon, France and Newtown, PA, USA, (2) University of California at Davis, CA, USA
- P58 bis** Validation of a multi-atlas segmentation technique for the quantification of hippocampal volume – Application to ADNI I
J.Schaerer (1), L.Bracoud (1), F.Roche (1), H.J.Yu (1), B.Belaroussi (1), D.Xu (1), C.Pachai (1), C.DeCarli (2) and the Alzheimer's Disease Neuroimaging Initiative
 (1) BioClinica Inc., Lyon, France and Newtown, PA, USA, (2) University of California at Davis, CA, USA

Biomarkers/Imaging in AD clinical trials - *Continued*

- P59** Changes of Biological Markers and Brain PET Imaging and Clinical Effects of Memantine for Patients with Moderate to Severe Alzheimer's Disease: a 24 Week Double-blind, Randomized, Placebo-Controlled Study
Shifu Xiao (1,2), Tao Wang (1,2), Qiu Huang (3), Kewei Chen (3,4), Eric Reiman (1,2,4)
 (1) Geriatric Psychiatry Department, Shanghai Mental Health Center, Shanghai JiaoTong University School of Medicine, Shanghai, China, (2) Alzheimer's Disease and Related Disorder Center, Shanghai JiaoTong University, Shanghai, China, (3) Med-X Institute, Shanghai Jiaotong University, Shanghai, China, (4) Banner Alzheimer's Institute and Banner Good Samaritan Medical Center, PET Center, Phoenix, AZ, USA
- P60** A new magnetic resonance-based approach to assessment of pathology in early Alzheimer's disease
Timothy James (1), Kristin James (1), Lance Farr (1), Gareth Thomas (1), James Rafferty (1), Michael Brady (2), David Chase (1)
 (1) Acuitas Medical Ltd., Swansea, Wales, UK, (2) Dept. of Engineering Science, University of Oxford, UK
- P61** Evaluation of brain fuel metabolism during normal aging and in MILD Alzheimer's disease (AD): comparison of 18F-Fluorodeoxyglucose with a novel PET ketone trace – Carbon –11 – Acetoacetate
Christian-Alexandre Castellano (1,2), Scott Nugent (1,2), Sébastien Tremblay (1,3), Mélanie Fortier (1), Nancy Paquet (3), Christian Bocti (4), Guy Lacombe (4), Éric Turcotte (3), Tamas Fulop (1,4), Stephen C Cunnane (1,2,4)
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- P62** A dual tracer PET and MRI approach to study deteriorating brain fuel metabolism during aging
Stephen C. Cunnane (1,2,3), Alexandre Courchesne-Loyer (1,2), Maggie Roy (1,2), Scott Nugent (1,2), Christian A. Castellano (1,2), Sébastien Tremblay (1,4), Éric Turcotte (4), Tamas Fulop (1,3)
 (1) Research Center on Aging, Sherbrooke, QC, Canada, (2) Department Physiology and Biophysics, Université de Sherbrooke, QC, Canada, (3) Department of Medicine, Université de Sherbrooke, QC, Canada, (4) Department of Radiobiology and Nuclear Medicine, Université de Sherbrooke, QC, Canada
- P63** An event- related potential index as a tool for the early diagnosis and drug development for Alzheimer's Disease
Vasileios Papaliagkas (1,2), Magda Tsolaki (2), Vasileios Kimiskidis (2)
 (1) Department of Neurology, Royal London Hospital, London, UK, (2) Third Neurological Clinic, Aristotle University of Thessaloniki, Greece

Nutrition and Alzheimer's Disease

- P64** Supporting synapse formation and function in Alzheimer's disease: mode of action of the specific nutrient combination Fortasyn™ Connect
John W Sijben, Martijn C de Wilde, Martine Groenendijk, Robert J Hageman, Patrick J Kamphuis, Laus M Broersen
 Nutricia Advanced Medical Nutrition, Danone Research, Centre for Specialised Nutrition, Wageningen, The Netherlands
- P65** Nutritional intervention with Fortasyn™ Connect: beneficial effects in experimental models of Alzheimer's pathology and functional decline
Nick van Wijk (1), Martijn C. de Wilde (1), Almar A.M. Kuipers (1), Martin Balvers (1), Martine Groenendijk (1), John W. Sijben (1), Patrick J. Kamphuis (1,2), Henna Koivisto (3,4), Heikki Tanila (3,4), Diane Jansen (5,6), Valerio Zerbi (5,6), Amanda J. Kiliaan (5,6), Laus M. Broersen (1)
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- P66** The Souvenaid Clinical Study Program for Alzheimer's disease
Philip Scheitens (1), Raj C Shah (2), David A Bennett (2), Rico L Wieggers (3), Tobias Hartmann (4,5), Hilka Soininen (6), Patrick JGH Kamphuis (3)
 (1) Department of Neurology, Alzheimer Center, VU University Medical Center, Amsterdam, The Netherlands, (2) Rush Alzheimer's Disease Center, Rush University Medical Center, Chicago, Illinois, United States, (3) Nutricia Advanced Medical Nutrition, Danone Research, Centre for Specialised Nutrition, Wageningen, The Netherlands, (4) Deutsches Institut für Demenz Prävention (DIDP), Neurodegeneration and Neurobiology, (5) Experimental Neurology, Homburg, Germany, (6) Department of Neurology, University of Eastern Finland and Kuopio University Hospital, Kuopio, Finland

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Nutrition and Alzheimer's Disease - *Continued*

- P67** *Update of the LipiDiDiet study : a study on the effects of the medical food Souvenaid in subjects with prodromal AD*
 Yvonne Freund-Levi (1), Pieter Jelle Visser (2,3), Mija Kivipelto (4), Patrick JGH Kamphuis (5), Rico L Wieggers (5), Tobias Hartmann (6,7), Hilka Soininen (8)
 (1) Department of NVS, Section of Clinical Geriatrics, Karolinska Institutet, Karolinska University Hospital, Huddinge, Sweden, (2) Department of Psychiatry and Neuropsychology, Alzheimer Center Limburg, University of Maastricht, The Netherlands, (3) Department of Neurology, Alzheimer Center, VU University Medical Center, Amsterdam, The Netherlands, (4) Aging Research Center, Karolinska Institutet and Stockholm Gerontology Research Center, Stockholm, Sweden, (5) Nutricia Advanced Medical Nutrition, Danone Research, Centre for Specialised Nutrition, Wageningen, The Netherlands, (6) Deutsches Institut für DemenzPrävention (DIP), Neurodegeneration and Neurobiology, (7) Experimental Neurology, Homburg, Germany, (8) Department of Neurology, University of Eastern Finland and Kuopio University Hospital, Kuopio, Finland
- P68** *Combined nutrient enrichment induces changes in synaptic protein expression in vitro*
 Paul J Savelkoul (1), Almar A Kuipers (1), Andrea Goudriaan (2), Robert J Hageman (1), John W Sijben (1), Guus AB Smit (2), Mark HG Verheijen (2), Patrick J Kamphuis (1,3), Laus M Broersen (1)
 (1) Nutricia Advanced Medical Nutrition, Danone Research, Centre for Specialised Nutrition, Wageningen, The Netherlands, (2) Dept. Molecular and Cellular Neurobiology, CNCR, VU University, Amsterdam, The Netherlands, (3) Utrecht Institute for Pharmaceutical Sciences (UIPS), University Utrecht, The Netherlands
- P69** *Alzheimer's Diet Modification: A Web-based Nutrition Tracking System For Patient Management and Outcomes Research*
 Richard S. Isaacson, Reza D. Khan, Christopher N. Ochner
 (1) Department of Neurology, University of Miami Miller School of Medicine, Miami USA, (2) Evolvix, NYC, USA, (3) Department of Psychology, Columbia University College of Physicians and Surgeons, NYC, USA
- P70** *Evidence on Diet Modification for Alzheimer's disease and Mild Cognitive Impairment*
 Christopher N. Ochner (1), Dulce Barrios (2), Clement Lee (2), Christine E. Greer (3), Richard S. Isaacson (3)
 (1)Columbia University, College of Physicians & Surgeons, NYC, USA, (2)Columbia University, Institute of Human Nutrition, NYC, USA, (3)University of Miami Miller School of Medicine, Miami, USA
- P71** *Souvenir II open-label extension study showed that Souvenaid was well-tolerated and suggests that memory performance continued to improve over 48 weeks*
 Philip Scheltens (1), Rafael Blesa (2), Marcel GM Olde Rikkert (3), Christine AF von Arnim (4), Anke Bongers (5), John Harrison (6), John Sijben (5), Elio Scarpini (7), Frans R Verhey (8), Maurits FJ Vandewoude (9), Rico L Wieggers (5), Bruno Vellas (10), Patrick JGH Kamphuis (5), on behalf of the Souvenir II open-label extension study group
 (1) Alzheimer Center, VU University Medical Center, Amsterdam, The Netherlands, (2) Hospital de la Sta Creu i St. Pau, Barcelona, Spain, (3) Alzheimer Centre Nijmegen, Radboud University Medical Centre Nijmegen, The Netherlands, (4) Department of Neurology, Ulm University, Ulm, Germany, (5) Nutricia Advanced Medical Nutrition, Danone Research, Centre for Specialised Nutrition, Wageningen, The Netherlands (6) Metis Cognition Ltd, Kilmington, UK & Imperial College, London, UK, (7) Ospedale Maggiore Policlinico, University of Milan, Milan, Italy, (8) Alzheimer Centre Limburg, Maastricht University Medical Centre, The Netherlands, (9) University Centre for Geriatrics, Antwerp Hospital Network (ZNA), Campus St Elisabeth, Antwerp, Belgium, (10) Gerontopole, INSERM U 1027, Toulouse, France

Miscellaneous in clinical trials

- P72** *Face-name associations memory training during non-invasive brain stimulation improves memory in Alzheimer's patients*
 Maria Cotelli (1), Rosa Manenti (1), Michela Petesi (1), Michela Brambilla (1), Sandra Rosini (1), Orazio Zanetti (1), Carlo Miniussi (1,3)
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- P73** *Physical exercise training in older adults diagnosed with mild to moderate dementia*
 Andreea Zamfirescu (1), Ana Capisizu (1), Mircea Slavila (2), Alexandru A. Capisizu (3), Aurel Romila (1)
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Miscellaneous in clinical trials - *Continued*

- P74** Cognitive impairment prevalence and correlations with subjective memory impairment: Findings from Brasov, Romania
Minerva Gurgu (1), Andreea Zamfirescu (2), Stroe Ana-Maria (3), Romila Aurel (1)
 (1) University of Medicine and Pharmacy "Carol Davila", Bucharest, Romania, Department of Geriatrics, Emergency Hospital- Brasov, Romania, (2) Clinic of Geriatrics, "Sf. Luca" Hospital, Bucharest, Romania (3) Department of Geriatrics, Hopitalier Bretagne, Vannes, France
- P75** Computerised cognitive testing can identify year by year declines in non-demented elderly aged 70 to 90 years
Keith A Wesnes (1,2), Brian K Saxby (3)
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- P76** The transition of cognitive decline from normal ageing to Mild Cognitive Impairment and Alzheimer's disease.
Keith Wesnes (1,2), William Lenderking (3)
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- P77** A cognitive task sensitive to dentate gyrus activity which has implications for assessing neurogenesis status in normal and pathological ageing
Keith A. Wesnes (1,2)
 (1) Bracket Global, Goring on Thames, United Kingdom, (2) Swinburne University, Melbourne, Australia
- P78** Use of the doll with Dementia-affected patients with behavior disturbances in a nursing home for Alzheimer patients.
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- P79** Therapeutic Virtual Train with Dementia-affected patients with behaviour disturbances in a nursing home for Alzheimer patients
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- P80** Assessment of an automated televigilance system on serious falls prevention in a dementia specialized care unit: the URCC
Isabelle Saulnier (1), Florent Lachal (1), Achille Tchalla (1), Justine Trimouillas (1), Florence Gourdeau-Nauché (2), Laurence Bernard-Bourzeix (1), Sophie Peyrichou (1), Sophie Fortuné (2), Thierry Dantoine (1)
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- P81** Robot assisted cognitive training can change the brain in the elderly: A single blind, randomized controlled trial of clinical efficacy
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- P82** Sepsis and Cognition
Catherine Widmann (1), Alexander Semmler (1,7), Torsten Okulla (1), Horst Urbach (3), Markus Kaiser (2,6), Guido Widman (4), Florian Mormann (4), Julia Weide (2), Klaus Fliessbach (4), Thomas Klockgether (1), Andreas Hoeft (2), Frank Jessen (5), Christian Putensen (2), Michael T. Heneka (2)
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SCIENTIFIC POSTERS

Miscellaneous in clinical trials - *Continued*

- P83** ADCS EDC
Gustavo A. Jimenez-Maggiola, Ronald G. Thomas, Phuoc Hong, Paul S. Aisen, Department of Neurosciences, UCSD, USA
- P84** High levels of CSF α -synuclein oligomers in Parkinson's disease with dementia and dementia with Lewy bodies but not in Alzheimer's disease
Omar M.A. El-Agnaf (1), Oskar Hansson (2,3,4), Sara Hall (2,3), Annika Öhrfelt (5), Shiji Varghese (1), Mohamed M. Qureshi (1), Abdulmonem Al-Hayani (6)
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- P85** Review of Safety of Spinal Catheter Placement for Intermittent Cerebrospinal Fluid Sampling in Clinical Trials on Healthy Volunteers
Steven Ramael (1), Luc Cavens (2), Anjana Dhar (1), Erika Wolters (1)
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- P86** A New 26-week, Double-blind, Randomized, Placebo-controlled, Study of AC-1204 (caprylic triglyceride) in Mild to Moderate Alzheimer's Disease: Presentation of Study Design
Rachelle Doody (1), James Galvin (2), Martin Farlow (3), Raj Shah (4), P. Murali Doraiswamy (5), Steven Ferris (2), Jean Zetlaoui (6), Samuel Henderson (7), Richard S. Isaacson (8)
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- P87** Prevention trials for Alzheimer's Disease in non-demented subjects. A review of studies that use biomarkers for inclusion or as outcome
Daniela Bertens (1), Pieter Jelle Visser (1,2), Philip Scheltens (1)
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- P88** Verbal Communication between Alzheimer's patients and caregivers: improving and enhancing communication
Renné P. Alegria (1,2), Cláudia C. Santana (2), Célia P. Gallo (2), Mirian G. Bolso (2), Cássio M. C. Bottino (2), Maria Inês Nogueira (1)
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Continuing Medical Education

Course Description :

Alzheimer's disease is one of the most important health challenges facing aging populations worldwide. The development of the next generation of Alzheimer's disease drugs is becoming essential to face up to this challenge. New pathways have been identified with biomarkers, facilitating novel trial designs for studies of tau-based therapies and other disease-modifying drugs including immunotherapy. However, methodological challenges continue to slow the development of specific new drug candidates. One of the objectives of the conference is to identify these hurdles and find ways to address them by bringing together world leaders in AD drug development to discuss solutions to the difficulties that have slowed the pace of progress, with a particular focus on clinical trial methodology.

Target Audience :

The target audience for CtaD2012 includes neurologists, psychiatrists and other clinicians and scientists involved in geriatric care, research, imaging and drug development for patients with Alzheimer's disease and other neurodegenerative disorders. Other healthcare professionals who may benefit from this activity are clinical research coordinators, nurses, speech therapists and other AD and dementia occupational therapists, psychologists and neuropsychologists.

Course Learning Objectives :

During this event essential learning objectives will be covered so that at the end of the conference each participant will be able to:

a) Translate the significance to the drug development process of each individual phase of clinical trials from Phase I to Phase IV.

- b) Articulate the involvement of neurotransmitters and cortical excitability in Alzheimer's disease pathology, and the implications for treatment.
- c) Implement in research studies: cognitive, clinical and biomarker measures that characterize the progression through the asymptomatic, prodromal and dementia phases of AD.
- d) Interpret back valuable information on new upcoming molecules in order to educate fellow physicians and propose alternatives to patients.
- e) List the novel methodologies or biomarkers essential in identifying predementia patients most at risk of developing AD.
- f) List study findings, including safety, biomarker and clinical findings, presented at the conference underlying progress in understanding the amyloid hypothesis.
- g) Assess different stages of AD, from the pre-symptomatic stages to severe dementia, to better plan and design future clinical trials
- h) Describe new composite outcome scores that optimize the power for measuring clinical disease progression for trials in a MCI and pre-MCI populations
- i) Describe and interpret sensitive biochemical (e.g. CSF A-beta, tau and phospho tau levels) and neurophysiological (e.g. QEEG, ERP) biomarkers of early AD detection.
- j) Interpret ERP data in AD clinical trials and drug development for cohort selection and monitoring disease progression.
- k) Identify and evaluate new methods for optimally scoring each patient's test item responses and overall test performance, for determining the number of dimensions underlying the test's performance, and for adjusting for sample bias effects.
- l) Express appreciation for vertical integration and the impact it can have by improving knowledge transfer and drug development

Continuing Medical Education



Needs assessments :

CTAD 2012 Scientific committee identified several practice gaps in designing and conducting AD Clinical Trials and developed the program to address these gaps, namely :

1. The need to appreciate the full spectrum of AD from an asymptomatic stage through dementia
2. The need for learning how dementia can affect specific population subgroups
3. The need to elucidate risk factors for AD and other dementias
4. The need to identify and avoid methodological errors in the design of multicenter and international clinical trials
5. The need for improved measurement of cognitive deficits
6. The need to keep up with scientific advances regarding biomarkers of AD pathologies.

Beyond these global practice gaps addressed throughout the conference other essential learning objectives will be covered so that at the end of the conference each participant will be able to understand :

- The importance of each individual phase of clinical trials from Phase I to Phase IV
- The relevance of neurotransmitters and cortical excitability in Alzheimer's Disease pathology
- The cognitive, clinical and biomarker measures that characterize the progression through the asymptomatic, prodromal and dementia phases of AD.

Accreditation Statement :

The University of California, San Diego School of Medicine is accredited by the Accreditation Council for Continuing Medical Education to provide continuing medical education for physicians.

Credit Designation Statement :

The University of California, San Diego School of Medicine designates this live activity for a maximum of **20.25 AMA PRA Category 1 Credits™**. Physicians should claim only the credit commensurate with the extent of their participation in the activity.

Cultural and Linguistic Competency Statement :

California Assembly Bill 1195 requires continuing medical education activities with patient care components to include curriculum in the subjects of cultural and linguistic competency. It is the intent of the bill, which went into effect on July 1, 2006, to encourage physicians and surgeons, CME providers in the state of California, and the Accreditation Council for Continuing Medical Education to meet the cultural and linguistic concerns of a diverse patient population through appropriate professional development. The planners, speakers and authors of this CME activity have been encouraged to address issues relevant in their topic area. In addition, a variety of resources are available that address cultural and linguistic competency, some of which may be included in your syllabus or handout materials. Additional resources and information about AB1195 can be found on our website at <http://cme.ucsd.edu>.

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