

## Wednesday, 23 September

Olympic Hall 1	Olympic Hall 2
	<b>08:30-09:00: Welcome – Opening ceremony</b> Theodoros Sklaviadis, Congress Chairman Jean-Philippe Deslys, Neuroprion Coordinator
	<b>9:00-9:30 Plenary talk</b> <b>Chair:</b> Theodoros Sklaviadis, Jean-Philippe Deslys Charles Weissmann, Jiali Li, Shawn Browning, Sukhvir P. Mahal, Anja Oelschlegel Department of Infectology, Scripps Florida, USA
	<b>Heterogeneity of prion populations</b>
<b>9:40-11:00: 1. Protein Misfolding</b> <b>Chair:</b> Claudio Soto, Umberto Agrimi	<b>9:40-10:55: 2. Diagnostics, Therapeutics &amp; Decontamination</b> <b>Chair:</b> Detlev Riesner, Hans Kretzschmar
<b>9:40-10:00</b> Tamar Canello <sup>1</sup> , Silvia Lisa <sup>2</sup> , Jakob Moskovitz <sup>3</sup> , Maria Gasset <sup>2</sup> , Ruth Gabizon <sup>1</sup> <sup>1</sup> Hadassah University Hospital, Israel; <sup>2</sup> Instituto de Química-Física "ROCASOLANO", Spain; <sup>3</sup> University of Kansas, USA <b>Oxidation of Met residues on PrP: An early event in prion formation and disease manifestation</b>	<b>09:40-10:00</b> <u>Inga Zerr</u> National TSE Reference Center, Göttingen, Germany <b>Clinical diagnosis of CJD: are there predictors of treatment response?</b>
<b>10:00-10:20</b> Agnieszka Salwierz <sup>1</sup> , <u>Eva Birkmann</u> <sup>1,2</sup> , Detlev Riesner <sup>1</sup> Heinrich-Heine Universität Düsseldorf, Germany; <sup>2</sup> Forschungszentrum Juelich <b>Studies on the interaction of membrane-bound PrP<sup>C</sup> with PrP<sup>Sc</sup></b>	<b>10:00-10:20</b> <u>Alexander Peden</u> <sup>1</sup> , Graham Fairfoul <sup>1</sup> , Suzanne Lowrie <sup>1</sup> , Linda McCardle <sup>1</sup> , Mark Head <sup>1</sup> , Seth Love <sup>2</sup> , Hester Ward <sup>1</sup> , Simon Cousens <sup>3</sup> , David Keeling <sup>4</sup> , Carolyn Millar <sup>5</sup> , FGH Hill <sup>6</sup> , James Ironside <sup>1</sup> <sup>1</sup> University of Edinburgh, UK; <sup>2</sup> Frenchay Hospital, Bristol, UK; <sup>3</sup> London School of Hygiene and Tropical Medicine, UK; <sup>4</sup> Churchill Hospital, Oxford, UK; <sup>5</sup> Imperial College London, UK; <sup>6</sup> Birmingham Children's Hospital, Birmingham, UK <b>vCJD infection in an asymptomatic UK haemophilic patient</b>
<b>10:20-10:40</b> <u>María Gasset</u> IQFR-CSIC, Spain <b>Structural intolerance of PrP α-fold for Helix-3 methionine oxidation: the link of theory and experiment</b>	<b>10:20-10:40</b> <u>Detlev Riesner</u> Institut für Physikalische Biologie, Heinrich-Heine-Universität Düsseldorf, Germany <b>Detection of prion particles in body fluids of humans and animals</b>
<b>10:40-11:00</b> <u>Will Guest</u> <sup>1</sup> , Li Li <sup>1</sup> , Olivier Julien <sup>2</sup> , Subhrangsu Chatterjee <sup>2</sup> , Brian Sykes <sup>2</sup> , Wenquan Zou <sup>3</sup> , Steven Plotkin <sup>1</sup> , Neil Cashman <sup>1</sup> <sup>1</sup> University of British Columbia, Canada, <sup>2</sup> University of Alberta, Canada, <sup>3</sup> Case Western Reserve University, USA <b>Partial unfolding of the prion protein: early steps on the path to misfolding</b>	<b>10:40-10:55</b> <u>Linda A. Terry</u> , Laurence Howells, Jeremy Hawthorn, Sally Everest, Sarah Jo Moore, Jane C. Edwards Veterinary Laboratories Agency, UK <b>Detection of prions in blood leucocytes</b>
<b>11:00-11:30 Coffee break</b>	
<b>11:30-12:50: 1. Protein Misfolding</b> <b>Chair:</b> Jesús Requena, Ruth Gabizon	<b>11:30-12:45: 2. Diagnostics, Therapeutics &amp; Decontamination</b> <b>Chair:</b> Maurizio Pocchiari, Inga Zerr
<b>11:30-11:50</b> <u>Federico Benetti</u> <sup>1</sup> , Heinz Amenitsch <sup>2,3</sup> , Matthijn Vos <sup>4</sup> , Adriana Ramos <sup>5</sup> , Peter Peters <sup>6</sup> , Giuseppe Legname <sup>1,3</sup> , Jesús Requena <sup>5</sup> <sup>1</sup> Scuola Internazionale Superiore di Studi Avanzati, Italy; <sup>2</sup> Austrian Academy of Sciences, Austria; <sup>3</sup> ELETTRA Laboratory, Sincrotrone Trieste S.C.p.A, Italy; <sup>4</sup> The FEI Company, Netherlands; <sup>5</sup> University of Santiago de Compostela, Spain; <sup>6</sup> Netherlands Cancer Institute, Netherlands <b>SAXS Analysis of PrP<sup>Sc</sup> and oligomeric recombinant mouse PrP(89-230)</b>	<b>11:30-11:45</b> <u>Chrysanthi Berberidou</u> , Konstantinos Xanthopoulos, Eleni Paulidou, Eleni Polyzoidou, Thanasis Lourbopoulos, Ioannis Poulos, Theodoros Sklaviadis Aristotle University of Thessaloniki, Greece <b>Decontamination of prion infected metal surfaces employing the photo-Fenton reagent</b>
	<b>11:45-12:00</b> <u>Silvio Notari</u> <sup>1</sup> , Liuting Qing <sup>1</sup> , Ayuna Dagdanova <sup>1</sup> , Sergei Ilchenko <sup>1</sup> , Mark E. Obrenovich <sup>1</sup> , Wen-Quan Zou <sup>1</sup> , Maurizio Pocchiari <sup>2</sup> , Pierluigi Gambetti <sup>1</sup> , Qingzhong Kong <sup>1</sup> , Shu G. Chen <sup>1</sup> <sup>1</sup> Case Western Reserve University, USA; <sup>2</sup> Istituto Superiore di Sanità, Italy <b>Human urine and PrP</b>

<p><b>11:50-12:10</b>  <u>Nathalie Daude</u><sup>1</sup>, Vivian Ng<sup>2</sup>, Joel Watts<sup>3</sup>, Sacha Genovesi<sup>1</sup>, Serene Wohlgemuth<sup>1</sup>, Jean-Paul Glaves<sup>1</sup>, Howard Young<sup>1</sup>, Gerold Schmitt-Ulms<sup>2</sup>, Joanne Mclaurin<sup>2</sup>, Paul E. Fraser<sup>2</sup>, David Westaway<sup>1</sup>  <sup>1</sup>University of Alberta, Canada; <sup>2</sup>University of Toronto, Canada; <sup>3</sup>University of San Francisco, USA  <b>The PrP-like Shadoo protein converts to amyloid at neutral pH</b></p>	<p><b>12:00-12:15</b>  <u>Kazuo Kuwata</u>, Tsutomu Kimura, Yuji O. Kamatari, Junji Hosokawa-Muto, Keiichi Yamaguchi, Takeshi Ishikawa, Takakazu Ishikura, Norifumi Yamamoto, Yumiko Okuda  Center for Emerging Infectious Diseases, Gifu University, Japan  <b>Rational design of anti-prion compounds targeting the PrP<sup>C</sup> characteristic sites</b></p>
<p><b>12:10-12:30</b>  <u>Human Rezaei</u><sup>1</sup>, Stephanie Prigent<sup>1</sup>, Nesrine Chakroun<sup>2</sup>, Celine Chapuis<sup>1</sup>, Cecile Dreiss<sup>2</sup>, Franca Fraternali<sup>2</sup>  <sup>1</sup>INRA, France; <sup>2</sup>King's College, UK  <b>Conformational dynamic of the PrP minimal domain involved in the oligomerization process</b></p>	<p><b>12:15-12:30</b>  <u>Andreas Mueller-Schiffmann</u><sup>1</sup>, Max Michel<sup>1</sup>, Benjamin Petsch<sup>2</sup>, Lothar Stitz<sup>2</sup>, Carsten Korth<sup>1</sup>  <sup>1</sup>University of Duesseldorf, Germany; <sup>2</sup>Friedrich Löffler Institute, Tuebingen, Germany  <b>A PrP<sup>Sc</sup> specific peptide derived from an antibody against PrP<sup>C</sup>/<sup>Sc</sup></b></p>
<p><b>12:30-12:50</b>  <u>Fabiana Caetano</u><sup>1,4</sup>, Iaci Soares<sup>1</sup>, Grace Pereira<sup>1</sup>, Marilene Lopes<sup>2</sup>, Nicolle Queiroz<sup>2</sup>, Stephen Ferguson<sup>3</sup>, Vania Prado<sup>1,3</sup>, Vilma Martins<sup>2</sup>, Marco Prado<sup>1,3</sup>  <sup>1</sup>Universidade Federal de Minas Gerais, Brazil; <sup>2</sup>Ludwig Institute for Cancer Research, Sao Paulo, Brazil; <sup>3</sup>University of Western Ontario, Canada  <b>Nuclear trafficking of ST11, a prion binding protein, induced by SUMOylation</b></p>	<p><b>12:30-12:45</b>  <u>Adriana Gielbert</u>, Mark Arnold, Peter Griffiths, John Spiropoulos, Maurice Sauer  Veterinary Laboratories Agency, UK  <b>N-Terminal amino acid profiling (N-TAAP) shows unexpected and distinct differences between PrPres from atypical and classical scrapie</b></p>
<b>13:00-14:20: Lunch (Meliton Hotel) / Poster Session (Olympic Hall 3)</b>	
	<p><b>14:30-15:00 Plenary talk</b>  Chair: Robert Will, Nikolaos Taskos</p>
	<p><u>Thomas Wisniewski</u>  New York University School of Medicine, USA  <b>Therapeutic Approaches for Alzheimer's and Prion Diseases</b></p>
	<p><b>15:10-16:30: 3. Other neurodegenerative proteinopathies</b>  Chair: Robert Will, Nikolaos Taskos</p>
	<p><b>15:10-15:30</b>  <u>Konrad Beyreuther</u>  Network Aging Research and ZMBH; University Heidelberg, Germany  <b>Novel therapeutic approaches in Alzheimer's disease</b></p>
	<p><b>15:30-15:50</b>  <u>Fabrizio Tagliavini</u><sup>1</sup>, Marcella Catania<sup>1</sup>, Michela Morbin<sup>1</sup>, Marco Gobbi<sup>2</sup>, Laura Colombo<sup>2</sup>, Antonio Bastone<sup>2</sup>, Laura Cantù<sup>3</sup>, Efrat Levy<sup>4</sup>, Maria Salmona<sup>2</sup>, Giuseppe Di Fede<sup>1</sup>  <sup>1</sup>"Carlo Besta" Neurological Institute, Milano, I; <sup>2</sup>"Mario Negri" Institute for Pharmacological Research, Milano, I; <sup>3</sup>University of Milan, Milano, I; <sup>4</sup>"Nathan S. Kline" Institute, Orangeburg, NY, USA  <b>Good protein, bad protein: A new Aβ variant can be both</b></p>
	<p><b>15:50-16:10</b>  <u>Magdalini Polymenidou</u><sup>1</sup>, Clotilde Lagier-Tourenne<sup>1</sup>, Tiffany Liang<sup>2</sup>, Gene Yeo<sup>2</sup> Don W. Cleveland<sup>1,2</sup>  <sup>1</sup>Ludwig Institute for Cancer Research, University of California in San Diego, USA; <sup>2</sup>Cellular and Molecular Medicine, University of California in San Diego, USA  <b>Role of RNA processing in the pathogenesis of amyotrophic lateral sclerosis</b></p>
<p><b>16:10-16:30</b>  <u>Hermann M. Schätzl</u>  Institute of Virology, Technische Universität München, Munich, Germany  <b>Autophagy and its role in neurodegenerative diseases</b></p>	
<b>16:30-17:00: Coffee break</b>	

<p><b>17:00-18:20: 4. Transmission &amp; Pathogenesis</b>  <b>Chair: Jean Manson, Eleni Kaldrymidou</b></p>	<p><b>17:00-18:20: 5. Basic Mechanisms of Neurodegeneration &amp; Pathology</b>  <b>Chair: Jan Langeveld, Edward Hoover</b></p>
<p><b>17:00-17:20</b>  <u>Jean Manson</u>  University of Edinburgh, UK  <b>TSE strains and their transmission within and between animals</b></p>	<p><b>17:00-17:20</b>  <u>Herbert Budka</u>  Institute of Neurology, Medical University Vienna, Austria  <b>Molecular neuropathology of TSEs</b></p>
<p><b>17:20-17:40</b>  <u>Olivier Andreoletti</u><sup>1</sup>, Nathalie Morel<sup>2</sup>, Caroline Lacroux<sup>1</sup>, Stéphanie Simon<sup>2</sup>, Séverine Lugan<sup>1</sup>, Jean-Louis Weisbecker<sup>3</sup>, Pierrette Costes<sup>1</sup>, Fabien Corbiere<sup>1</sup>, Jacques Grassi<sup>1</sup>, François Schelcher<sup>1</sup>  <sup>1</sup>UMR INRA ENVT 1225 Interactions Hôte agent pathogène, France; <sup>2</sup>CEA, Service de Pharmacologie et d'Immunoanalyse, IBItec-S, DSV, CEA/Saclay, France; <sup>3</sup>INRA Domaine de Langlade, France  <b>TSE genetic resistance and allelic interference in sheep</b></p>	<p><b>17:20-17:40</b>  <u>Martin Jeffrey</u><sup>1</sup>, Bruce Chesebro<sup>2</sup>, Brent Race<sup>2</sup>, Gillian McGovern<sup>1</sup>  <sup>1</sup>Veterinary Laboratories Agency, UK; <sup>2</sup>Rocky Mountain Laboratories, USA  <b>Patterns of abnormal PrP drainage and amyloid formation in scrapie infected mice expressing GPI-anchorless PrP</b></p>
<p><b>17:40-18:00</b>  <u>Edgar Holznel</u><sup>1</sup>, Walter Schulz-Schaeffer<sup>2</sup>, Barbara Yutzky<sup>1</sup>, Gerhard Hunsmann<sup>3</sup>, Johannes Loewer<sup>1</sup>  <sup>1</sup>Paul-Ehrlich-Institut, Federal Institute for Sera and Vaccines, Germany; <sup>2</sup>Department of Neuropathology, Georg-August University, Göttingen, Germany, <sup>3</sup>Department of Virology and Immunology, German Primate Centre, Göttingen, Germany  <b>Spread of BSE prions in cynomolgus monkeys (<i>Macaca fascicularis</i>) after oral transmission</b></p>	<p><b>17:40-18:00</b>  <u>Pedro Fernandez-Funez</u><sup>1</sup>, Sergio Casas-Tinto<sup>1</sup>, Yan Zhang<sup>1</sup>, Wen-Quan Zou<sup>2</sup>, Diego Rincon-Limas<sup>1</sup>  <sup>1</sup>University of Texas Medical Branch, USA, <sup>2</sup>Case Western Reserve University, USA  <b>Sequence-dependent pathobiology of prion proteins: lessons from rabbits, mice and hamsters</b></p>
<p><b>18:00-18:20</b>  <u>Anne Buschmann</u><sup>1</sup>, Ute Ziegler<sup>1</sup>, Leila McIntyre<sup>2</sup>, Markus Keller<sup>1</sup>, Ron Rogers<sup>3</sup>, Bob Hills<sup>3</sup>, Martin H. Groschup<sup>1</sup>  <sup>1</sup>Friedrich-Loeffler-Institut, INEID, Germany; <sup>2</sup>Faculty of Veterinary Medicine, University of Calgary, Canada; <sup>3</sup>Health Canada, Ottawa, Canada  <b>PrP<sup>Sc</sup> distribution pattern in cattle experimentally challenged with H-type and L-type atypical BSE</b></p>	<p><b>18:00-18:20</b>  Sabrina Cronier<sup>1</sup>, <u>Julie Carimalo</u><sup>2</sup>, Brigitte Schaeffer<sup>3</sup>, Marie-Christine Miquel<sup>4</sup>, Hubert Laude<sup>1</sup>, Jean-Michel Peyrin<sup>5</sup>  <sup>1</sup>INRA U892 VIM, France; <sup>2</sup>Reference Center for TSE Surveillance UMG, Germany; <sup>3</sup>INRA MIA, France; <sup>4</sup>CNRS UMR 5241 UPS, France; <sup>5</sup>CNRS UMR 7102 UPMC, France  <b>Endogenous PrP conversion is required for prion-induced neuritic alterations and neuronal death</b></p>
<p><b>18:20-18:50</b>  Talks by two young researchers from the Neuroprion network, selected after training in scientific communication  <b>Chair: Jean Philippe Deslys</b></p>	<p><b>18:20-18:50</b>  Talks by two young researchers from the Neuroprion network, selected after training in scientific communication  <b>Chair: Jean Philippe Deslys</b></p>
<p><b>18:50-20:30: Poster Party (Olympic Hall 3)</b></p>	

Thursday, 24 September

Olympic Hall 1	Olympic Hall 2
	<p><b>Plenary talk</b>  <b>Chair: Juan Maria Torres, Stavros Balogiannis</b></p>
	<p><b>9:00-9:30</b>  <u>Laura Manuelidis</u>            Yale University Medical School, USA</p>
	<p><b>What agent strains tell us</b></p>
<p><b>10:00-11:00: 5. Basic mechanisms of neurodegeneration &amp; Pathology</b>  <b>Chair: Herbert Budka, Maria Gasset</b></p>	<p><b>9:30-11:00: 4. Transmission &amp; Pathogenesis</b>  <b>Chair: Juan Maria Torres, Stavros Balogiannis</b></p>
<p><b>10:00-10:20</b>  <u>Benoit Schneider</u><sup>1</sup>, Damien Loubet<sup>1</sup>, Elodie Pradines<sup>1</sup>, Yannick Bailly<sup>2</sup>, Jean-Michel Peyrin<sup>3</sup>, Hubert Laude<sup>3</sup>, Catherine Vidal<sup>1</sup>, Jean-Marie Launay<sup>1</sup>, Sophie Mouillet – Richard<sup>1</sup>, Odile Kellermann<sup>1</sup>  <sup>1</sup>INSERM U747 - Université Paris Descartes, France; <sup>2</sup>CNRS UPR 3212, France; <sup>3</sup>INRA, France</p> <p><b>Prion replication sensitized neurons to pro-inflammatory cytokines</b></p>	<p><b>9:30-10:00</b>  <u>Edward A. Hoover</u><sup>1</sup>, Nicholas J. Haley<sup>1</sup>, Candace K. Mathiason<sup>1</sup>, Nathaniel D. Denkers<sup>1</sup>, Davis M. Seelig<sup>1</sup>, and Glenn C. Telling<sup>2</sup>  <sup>1</sup>Colorado State University, USA; <sup>2</sup>University of Kentucky, Lexington, USA</p> <p><b>Recent results on the transmission, detection, and pathogenesis of chronic wasting disease</b></p>
<p><b>10:20-10:40</b>            Anna Majer<sup>1</sup>, Sarah Medina<sup>2</sup>, Reuben Saba<sup>1</sup>, Debra Parchaliuk<sup>2</sup>, Kathy Frost<sup>2</sup>, Catherine Robertson<sup>2</sup>, <u>Stephanie Booth</u><sup>2</sup>  <sup>1</sup>University of Manitoba, Canada; <sup>2</sup>Public Health Agency of Canada, Canada</p> <p><b>The investigation of dysregulated microRNAs in prion – induced neurodegeneration</b></p>	<p><b>10:00-10:20</b>  <u>Sandra McCutcheon</u><sup>1</sup>, Anthony Richard Alejo Blanco<sup>1</sup>, Christopher de Wolf<sup>1</sup>, Boon Chin Tan<sup>1</sup>, Nora Hunter<sup>1</sup>, Valerie Hornsey<sup>2</sup>, Christopher Prowse<sup>2</sup>, Marc Turner<sup>2</sup>, Martin H Groschup<sup>3</sup>, Dietmar Becher<sup>4</sup>, Fiona Houston<sup>5</sup>, Jean C Manson<sup>1</sup>  <sup>1</sup>The Roslin Institute and R (D) SVS, University of Edinburgh, UK; <sup>2</sup>Scottish National Blood Transfusion Service, UK; <sup>3</sup>FLI-Federal Research Institute for Animal Health, Germany; <sup>4</sup>Micromun, Germany; <sup>5</sup>University of Glasgow, UK</p> <p><b>All separated components, prepared from BSE-infected sheep blood, are infectious upon transfusion</b></p>
<p><b>10:40-11:00</b>  <u>Mohammed Moudjou</u>, Michel Dron, Jérôme Chapuis, Julie Bernard, Muhammad K. F. Salamat, Sabrina Cronier, Christelle Langevin, Hubert Laude            INRA, France</p> <p><b>Endogenous proteolytic cleavage of disease-associated prion protein to produce C2 fragments is strongly cell- and tissue-dependent</b></p>	<p><b>10:20-10:40</b>  <u>Lorenzo Gonzalez</u><sup>1</sup>, Silvia Siso<sup>1</sup>, Stuart Martin<sup>1</sup>, Steve A.C. Hawkins<sup>1</sup>, Wilfred Goldmann<sup>2</sup>, Timm Konold<sup>1</sup>, James Hope<sup>1</sup>, Martin Jeffrey<sup>1</sup>  <sup>1</sup>Veterinary Laboratories Agency, UK; <sup>2</sup>Roslin Institute, UK</p> <p><b>Pathogenesis of scrapie in goats: modulation by host PRNP genotype and effect of coexistent conditions</b></p>
	<p><b>10:40-11:00</b>            Xiangmei Zhou<sup>1</sup>, Haiyun Zhou<sup>1</sup>, Mohammed Kouadir<sup>1</sup>, Zhongqiu Zhang<sup>2</sup>, Lifeng Yang<sup>1</sup>, <u>Deming Zhao</u><sup>1</sup>  <sup>1</sup>China Agricultural University; <sup>2</sup>China Animal Disease Control Center, China</p> <p><b>Induction of macrophage migration by neurotoxic prion protein fragment</b></p>
<b>11:00-11:30 Coffee break</b>	
<p><b>11:30-12:50: 6. Functions &amp; Cell biology of PrP</b>  <b>Chair: Olivier Andreoletti, Magda Tsolaki</b></p>	<p><b>11:30-13:00: 7. Genetics</b>  <b>Chair: Fabrizio Tagliavini, Pierluigi Gambetti</b></p>
<p><b>11:30-11:50</b>  <u>Cathryn Haigh</u><sup>1</sup>, Amelia McGlade<sup>1</sup>, Simon Drew<sup>1</sup>, Martin Boland<sup>1</sup>, Colin Masters<sup>2</sup>, Kevin Barnham<sup>1</sup>, Victoria Lawson<sup>1</sup>, Steven Collins<sup>1</sup>  <sup>1</sup>The University of Melbourne, Australia; <sup>2</sup>The Mental Health Research Institute, Australia</p> <p><b>Synergy of polybasic structure and copper co-ordination in the PrP23-89-mediated stress protection response</b></p>	<p><b>11:30-12:00</b>  <u>John Collinge</u>            MRC Prion Unit and Department of Neurodegenerative Disease, UCL Institute of Neurology, National Hospital for Neurology and Neurosurgery, Queen Square, London</p> <p><b>Understanding genetic susceptibility to human prion diseases</b></p>
<p><b>11:50-12:10</b>  <u>Gerold Schmitt-Ulms</u><sup>1</sup>, Joel Watts<sup>2</sup>, Sepehr Ehsani<sup>1</sup>, Hairu Huo<sup>1</sup>, Yu Bai<sup>3</sup>, Holger Wille<sup>2</sup>, David Westaway<sup>4</sup>  <sup>1</sup>University of Toronto, Canada; <sup>2</sup>University of California San Francisco, USA; <sup>3</sup>Peking University, China; <sup>4</sup>University of Alberta, Canada</p> <p><b>Family reunion - encounter of molecular cousins uncovers founder and ancient function of prion gene family</b></p>	<p><b>12:00-12:30</b>  <u>David Westaway</u><sup>1,2,3</sup>, Nathalie Daude<sup>1</sup>, Sacha Genovesi<sup>1</sup>, Inyoul Lee<sup>4</sup>, Jing Yang<sup>1</sup>, Rebecca Young<sup>5</sup>, and George A. Carlson<sup>5</sup>  <sup>1</sup>Centre for Prions and Protein Folding Diseases, University of Alberta; <sup>2</sup>Division of Neurology, Faculty of Medicine, University of Alberta; <sup>3</sup>Department of Biochemistry, University of Alberta; <sup>4</sup>Institute for Systems Biology, University of Washington; <sup>5</sup>McLaughlin Research Institute, Great Falls Montana</p> <p><b>Properties of the Shadoo Protein in Health and Prion Disease</b></p>

<p><b>12:10-12:30</b>  <u>Michel Dron</u><sup>1</sup>, Françoise Dandoy-Dron<sup>2</sup>, Muhammad Khalid Farooq Salamat<sup>1</sup>, Hubert Laude<sup>1</sup>  <sup>1</sup>INRA, France; <sup>2</sup>CNRS, France</p> <p><b>Proteasome inhibitors induce accumulation of a detergent-insoluble PrP<sup>26K</sup> and formation of PrP<sup>Sc</sup> aggregates in prion-infected CAD cells</b></p>	<p><b>12:30-12:45</b>  <u>Simon Mead</u>, Jerome Whitfield, Mark Poulter, Paresh Shah, James Uphill, Tracy Campbell, Huda Al-Dujaily, Holger Hummerich, Jon Beck, Charles A Mein, Claudio Verzilli, John Whittaker, Michael Alpers, John Collinge  Medical Research Council, UK</p> <p><b>A novel protective prion protein variant co-localises with kuru exposure</b></p>
<p><b>12:30-12:50</b>  <u>Rafael Mariante</u>, Alberto Nóbrega, Maria Bellio, Rafael Liden  Federal University of Rio de Janeiro</p> <p><b>Neuroimmunoendocrine regulation of the prion protein in neutrophils</b></p>	<p><b>12:45-13:00</b>  <u>Jon Beck</u><sup>1</sup>, Diego Kaski<sup>1</sup>, Catherine Pennington<sup>2</sup>, Mark Poulter<sup>1</sup>, James Uphill<sup>1</sup>, Matthew Bishop<sup>2</sup>, Jackie Linehan<sup>1</sup>, Catherine O'Malley<sup>1</sup>, Sebastian Brandner<sup>1</sup>, Richard Knight<sup>3</sup>, John Collinge<sup>1</sup>, Simon Mead<sup>1</sup>  <sup>1</sup>MRC Prion Unit at the National Prion Clinic, UK; <sup>2</sup>National Hospital for Neurology and Neurosurgery, UK; <sup>3</sup>National CJD Surveillance Unit, UK</p> <p><b>Inherited prion disease with 4-octapeptide repeat insertion – disease requires the interaction of multiple genetic risk factors</b></p>
<b>13:00-14:20 Lunch (Meliton Hotel) / Poster session (Olympic Hall 3)</b>	
	<p><b>14:30-16:35: 8. IPFA/FABS/LFB: Blood safety and prions</b>  Chair: Paul Brown</p>
	<p><b>14:30-14:55</b>  <u>Robert G. Will</u>  National CJD Surveillance Unit, Edinburgh, UK</p> <p><b>Variant CJD and plasma products</b></p>
	<p><b>14:55-15:20</b>  <u>Marc Turner</u>  Scottish National Blood Transfusion Service and Department of Haematology, Royal Infirmary, Edinburgh, UK</p> <p><b>Blood safety: from screening tests to prion removal</b></p>
	<p><b>15:20-15:45</b>  Byron Caughey  Rocky Mountain Laboratories, NIAID, NIH, Hamilton, MT 59840</p> <p><b>Structural analyses and detection of prion proteins: from rapid diagnosis to therapeutic development</b></p>
	<p><b>15:45-16:10</b>  <u>Claudio Soto</u>, Marcelo Barria, Baian Chen, Sense Gonzalez-Romero, Abhisek Mukherjee, Rodrigo Morales  University of Texas Medical School at Houston, USA</p> <p><b>Quantification of PrP<sup>Sc</sup> in different blood fractions, brain and peripheral tissues at distinct stages of prion disease</b></p>
<p><b>16:10-16:35</b>  <u>Conrad G. Brunk</u>  Department of Philosophy, University of Victoria, Canada</p> <p><b>Ethical implications of prion blood screening tests</b></p>	
<b>16:35-17:00 Coffee break</b>	
<p><b>17:00-18:20: 9. Epidemiology &amp; Risk assessment</b>  Chair: Michael Dawson, Nikolaos Papaioannou</p>	<p><b>17:00-18:35: 10. Natural &amp; experimental strains</b>  Chair: Hubert Laude, Penelope Papisavva Stylianou</p>
<p><b>17:00-17:20</b>  Alexandre Fediaevsky<sup>1</sup>, Cristiana Maurella<sup>2</sup>, Maria Nöremark<sup>3</sup>, Francesco Ingravalle<sup>2</sup>, Stefania Thorgeirsdottir<sup>4</sup>, Leonor Orge<sup>5</sup>, Renaud Poizat<sup>6</sup>, Maria Hautaniemi<sup>7</sup>, Barry Liam<sup>8</sup>, Didier Calavas<sup>9</sup>, <u>Giuseppe Ru</u><sup>2</sup>, Petter Hopp  <sup>1</sup>AFSSA-Lyon; INRA Clermont-Theix, France; <sup>2</sup>Istituto Zooprofilattico Sperimentale del Piemonte, Liguria e Valle d'Aosta, Italy; <sup>3</sup>National Veterinary Institute, Sweden; <sup>4</sup>University of Iceland, Iceland; <sup>5</sup>Laboratório Nacional de Investigação Veterinária, Portugal; <sup>6</sup>AFSCA, Portugal; <sup>7</sup>Finnish Food Safety Authority, Finland; <sup>8</sup>Department of Agriculture, Fisheries and Food, Ireland; <sup>9</sup>AFSSA-Lyon, France; <sup>10</sup>National Veterinary Institute, Norway</p> <p><b>No evidence of contagiousness of atypical scrapie: results from a 12-country European study</b></p>	<p><b>17:00-17:20</b>  <u>Vincent Béringue</u>, Hubert Laude.  INRA, UR 892, Virologie Immunologie Moléculaires, France</p> <p><b>Transmission of atypical bovine prions to mice transgenic for human prion protein</b></p> <p><b>17:20-17:35</b>  <u>David Colby</u>, Giuseppe Legname, Holger Wille, Ilia Baskakov, Stephen DeArmond, Stanley Prusiner  University of California, San Francisco, USA</p> <p><b>Generation of diverse synthetic prion strains capable of infecting mice that express full-length prion protein</b></p>

<p><b>17:20-17:40</b>  <u>Robert Somerville</u>, Karen Fernie, Allister Smith  University of Edinburgh, UK</p> <p><b>Survival and limited spread of TSE infectivity after burial for one year</b></p>	<p><b>17:35-17:50</b>  <u>Umberto Agrimi</u><sup>1</sup>, Michele Di Bari<sup>1</sup>, Claudia D'Agostino<sup>1</sup>, Elena Esposito<sup>1</sup>, Maria Mazza<sup>2</sup>, Simone Barocci<sup>3</sup>, Geraldina Riccardi<sup>1</sup>, Shimon Simson<sup>1</sup>, Paolo Frassanito<sup>1</sup>, Barbara Iulini<sup>2</sup>, Pier Luigi Acutis<sup>2</sup>, Cristina Casalone<sup>2</sup>, Hubert Laude<sup>4</sup>, Sylvie Benestad<sup>5</sup>, Gabriele Vaccari<sup>1</sup>, Romolo Nonno<sup>1</sup></p> <p><sup>1</sup>Istituto Superiore di Sanità, Italy; <sup>2</sup>Istituto Zooprofilattico Sperimentale del Piemonte, Liguria e Valle d'Aosta, Italy; <sup>3</sup>Istituto Zooprofilattico Sperimentale dell'Umbria e delle Marche, Italy, <sup>4</sup>INRA, France; <sup>5</sup>National Veterinary Institute, Norway</p> <p><b>Separation of prion strains from a mixture by discriminatory bioassay. The example of a natural case of mixed infection of classical scrapie and Nor98</b></p>
<p><b>17:40-18:00</b>  <u>Hong Yang</u>, Richard Forshee, Mark Walderhaug, Steven Anderson  US Food and Drug Administration, USA</p> <p><b>Updated risk assessment of variant Creutzfeldt-Jakob disease (vCJD) risks for recipients of plasma-derived blood clotting products in the US</b></p>	<p><b>17:50-18:05</b>  <u>Silvia Sisó</u><sup>1</sup>, Francesca Chianini<sup>2</sup>, Scott Hamilton<sup>2</sup>, Mark Dagleish<sup>2</sup>, Philip Steele<sup>2</sup>, Jeanie Finlayson<sup>2</sup>, Stuart Martin<sup>1</sup>, Martin Jeffrey<sup>1</sup>, Lorenzo González<sup>1</sup></p> <p><sup>1</sup>Veterinary Laboratories Agency- Lasswade, UK; <sup>2</sup>Moredun Research Institute, UK</p> <p><b>Significance of murine scrapie strains: ME7 changes phenotype when inoculated in sheep and then back into mice</b></p>
<p><b>18:00-18:20</b>  <u>Sylvie Benestad</u><sup>1</sup>, Torfinn Moldal<sup>1</sup>, Torkjel Bruheim<sup>2</sup>, Petter Hopp<sup>1</sup>, Bjørn Bratberg<sup>1</sup></p> <p><sup>1</sup>National Veterinary Institute, Oslo, Norway, <sup>2</sup>National Veterinary Institute, Trondheim, Norway</p> <p><b>Is Nor98 scrapie contagious? Results from Norwegian flock studies</b></p>	<p><b>18:05-18:20</b>  <u>Pierluigi Gambetti</u>  Case Western Reserve University, USA</p> <p><b>A novel human prion disease affecting subjects with the three prion protein codon 129 genotypes: Could it be the sporadic form of Gerstmann-Sträussler</b></p>
	<p><b>18:20-18:35</b>  <u>Romolo Nonno</u><sup>1</sup>, Michele Di Bari<sup>1</sup>, Laura Pirisinu<sup>1</sup>, Stefano Marcon<sup>1</sup>, Claudia D'Agostino<sup>1</sup>, Elena Esposito<sup>1</sup>, Paola Fazzi<sup>1</sup>, Shimon Simson<sup>1</sup>, Paolo Frassanito<sup>1</sup>, Cristina Casalone<sup>3</sup>, Franco Cardone<sup>2</sup>, Maurizio Pocchiari<sup>2</sup>, Gabriele Vaccari<sup>1</sup>, Umberto Agrimi<sup>1</sup></p> <p><sup>1</sup>Dept. SPVSA, Istituto Superiore di Sanità, Italy; <sup>2</sup>Dept. BCN, Istituto Superiore di Sanità, Italy; <sup>3</sup>Istituto Zooprofilattico del Piemonte, Liguria e Valle D'Aosta, Italy</p> <p><b>Biological typing of sporadic Creutzfeldt-Jakob disease isolates and comparison with animal prion isolates</b></p>
	<p><b>18:35-19:05: Plenary talk</b>  Chair: Paul Brown</p> <p><u>Lev G. Goldfarb</u>  Bethesda, Maryland, U.S.A.</p> <p><b>In memory of Dr Daniel Carleton Gajdusek (1923 - 2008)</b></p> <p><b>Announcement of poster prize winners</b>  <b>Announcement of the grants awarded for the 2009 call by the Alliance Biosecure Foundation</b></p> <p><b>19:10-19:40: Anne-Charlotte Panissié</b></p> <p><b>A journey through Greek mythology and history, the figure of Homer</b></p>
<p><b>20:30 Gala Dinner (Meliton Hotel)</b></p>	

Friday, 25 September

<b>Olympic Hall 2</b>
<b>8:15-9:30: 11. Hot topics</b> <b>Chair:</b> Hermann Schätzl, Vasilios Kokkas
<b>8:15-8:30</b> <b>Gültekin Tamgüney</b> <sup>1,2</sup> , Michael W. Miller <sup>3</sup> , Lisa L. Wolfe <sup>3</sup> , Tracey M. Sirochman <sup>3</sup> , David V. Glidden <sup>4</sup> , Christina Palmer <sup>1</sup> , Azucena Lemus <sup>5</sup> , Stephen J. DeArmond <sup>5</sup> , Stanley B. Prusiner <sup>1,2</sup> <sup>1</sup> Institute for Neurodegenerative Diseases, University of California, San Francisco, USA; <sup>2</sup> Department of Neurology, University of California, San Francisco, USA; <sup>3</sup> Colorado Division of Wildlife, Wildlife Research Center, Fort Collins, USA; <sup>4</sup> Department of Epidemiology and Biostatistics, University of California, San Francisco, USA; <sup>5</sup> Department of Pathology, University of California, San Francisco, USA
<b>Prions in feces of asymptomatic deer</b>
<b>8:30-8:45</b> <b>Emmanuel Comoy</b> <sup>1</sup> , Juergen Richt <sup>2</sup> , Valérie Durand <sup>1</sup> , Sophie Freire <sup>1</sup> , Evelyne Correia <sup>1</sup> , Amir Hamir <sup>2</sup> , Marie-Madeleine Ruchoux <sup>1</sup> , Paul Brown <sup>1</sup> , Jean-Philippe Deslys <sup>1</sup> <sup>1</sup> Atomic Energy Commission, France; <sup>2</sup> National Animal Disease Center, USA
<b>Transmission of bovine-passaged TME prion strain to macaque</b>
<b>8:45-9:00</b> <b>Silvia Suardi</b> <sup>1</sup> , Chiara Vimercati <sup>1</sup> , Fabio Moda <sup>1</sup> , Ruggerone Margherita <sup>1</sup> , Ilaria Campagnani <sup>1</sup> , Guerino Lombardi <sup>2</sup> , Daniela Gelmetti <sup>2</sup> , Martin H Groschup <sup>3</sup> , Anne Buschmann <sup>3</sup> , Cristina Casalone <sup>4</sup> , Maria Caramelli <sup>4</sup> , Salvatore Monaco <sup>5</sup> , Gianluigi Zanusso <sup>5</sup> , Fabrizio Tagliavini <sup>1</sup> <sup>1</sup> Carlo Besta" Neurological Institute, Italy; <sup>2</sup> IZS Brescia, Italy; <sup>3</sup> 3FLI Insel Riems, D, Germany; <sup>4</sup> CEA-IZS Torino, Italy; <sup>5</sup> University of Verona, Italy
<b>Infectivity in skeletal muscle of BASE-infected cattle</b>
<b>9:00-9:15</b> <b>Edward Málaga-Trillo</b> , Gonzalo P. Solis, Claudia A.O. Stuermer, Yvonne Schrock, Emily Sempou, Corinna Geiss Department of Biology, University of Konstanz, Germany
<b>The use of zebrafish in prion biology: lessons and perspectives</b>
<b>9:15-9:30</b> Steve Simoneau, Jean-Guy Fournier
<b>Small critical RNAs in the scrapie agent</b>
<b>9:30-10:20 Round table : Sustaining prion research: from past to future</b> <b>Chair:</b> Christian Patermann Jim Hope: Situation in UK Neil Cashman: Developing in Canada a full spectrum of research activities on prion Charles Weissmann: Prion research from Europe to US Christian Patermann: Synthesis: strengthening a durable integration of prion research
<b>10:20-10:35 Family session</b>
CJD International Support Alliance, Building a Global Prion Disease Support Community Florence Kranitz, President of the CJD Foundation Susane Solvyns, CJD Support Group Network, Australia; Co-Chair of the CJD International Support Alliance
<b>10:35-11:45: Poster prize winners presentations</b> <b>Chair:</b> Jean Philippe Deslys, David Westaway
<b>11:45-12:00 Closing ceremony / Next meeting announcement</b>