

## Poster Presentations List

### Poster Number:

- 1. The Role of MSWS Concerning The Improvement of Autopsy Rates of Prion Diseases**  
Shoken Aizawa, Mihara Memorial Hospital
- 2. Prion Protein Immunohistochemistry Using Automatic Immunostainer**  
Shinichi Aoyagi, Mihara Memorial Hospital
- 3. Synthesis and evaluation of radioiodinated flavonoid-related compounds as SPECT probes for imaging cerebral prion deposits**  
Takeshi Fuchigami, Nagasaki University
- 4. Thin-slice diffusion-weighted imaging and arterial spin labeling for the diagnosis of Creutzfeldt-Jakob disease**  
Koji Fujita, The University of Tokushima
- 5. Detection specificity of salivary BSE PrP<sup>Sc</sup> by PMCA**  
Shigeo Fukuda, Hokkaido Animal Research Center
- 6. Serial magnetic resonance imaging changes in a patient in the early stages of sporadic Creutzfeldt-Jacob disease with valine homozygosity at codon 129 of *PRNP***  
Fumiko Furukawa, Tokyo Medical and Dental University
- 7. A family of hereditary progressive dementia with five-octapeptide repeat insertion in prion protein gene : An unique case of genetic Creutzfeldt-Jakob disease**  
Shinsuke Hamada, Hokuyukai Neurological Hospital
- 8. Comparison of dura mater graft-associated Creutzfeldt-Jakob disease between Japan and other countries**  
Tsuyoshi Hamaguchi, Kanazawa University
- 9. Reaction of complement factors on scrapie-infected primary-cultured neurons temporarily increases permeability of plasma membrane**  
Rie Hasebe, Hokkaido University
- 10. Clinicopathological findings of an autopsy case of MM2-thalamic-type sporadic Creutzfeldt-Jakob disease**  
Yuichi Hayashi, Gifu University
- 11. Human prion diseases in Japan: a prospective surveillance from 1999**  
Masaki Hizume, Tokyo Medical and Dental University/ Saitama Prefecture Rehabilitation Center

- 12. Analysis of microglial activation state in brains of prion-infected mice**  
Motohiro Horiuchi, Hokkaido University
- 13. Neuropathological investigation of the olfactory bulb and olfactory tract in sporadic Creutzfeldt-Jakob disease**  
Yasushi Iwasaki, Aichi Medical University
- 14. Identification and characterization of multispecific monoclonal antibody G2 directed against chicken prion protein**  
Yuji O. Kamatari, Gifu University
- 15. Identification of the cryptic mitochondrial targeting sequence for PrP<sup>C</sup>/mitochondria-dependent neuronal cell death**  
Hiroki Kato, Tokyo Medical University
- 16. Establishment of transgenic mice expressing a human prion protein**  
Chi-Kyeong Kim, Korea Centers for Disease Control & Prevention
- 17. Revision of Sporadic Creutzfeldt-Jakob Disease with MV2 prions**  
Atsushi Kobayashi, Tohoku University
- 18. Mechanisms of anti-prion factors in yeast**  
Hiroshi Kurahashi, Tohoku University
- 19. Type I IFN signal pathway of host innate immune response regulates prion infection**  
Daisuke Ishibashi, Nagasaki University
- 20. The emergence of novel BSE prions by the serial transmission of H-type BSE in bovinized mice**  
Kentaro Masujin, National Institute of Animal Health
- 21. Transmissible properties of field scrapie isolates to GT1-7 cells**  
Kohtaro Miyazawa, National Institute of Animal Health
- 22. Linear epitopes on the surface of PrP<sup>res</sup> of sporadic Creutzfeldt-Jakob disease**  
Masanori Morita, Japan Blood Products Organization
- 23. FK506 reduces abnormal prion protein through the activation of autolysosomal degradation and prolongs survival in prion-infected mice**  
Takehiro Nakagaki, Nagasaki University
- 24. Prions impair post-Golgi trafficking of membrane proteins**  
Suehiro Sakaguchi, The University of Tokushima
- 25. Glycerol enhances the PrP<sup>res</sup> production via a PI3K signaling pathway in prion-infected neuroblastoma cells**  
Yuji Sakasegawa, Tohoku University

- 26. Prospective surveillance data of human prion disease in Chugoku and Shikoku regions in Japan**  
Kota Sato, Okayama University
- 27. Brain microglia were silent in fatal familial insomnia and G114V genetic CJD, but activated in sporadic CJD**  
Qi Shi, Chinese Center for Disease Control and Prevention
- 28. PrP octarepeats region determined the interaction with caveolin-1 and phosphorylation of caveolin-1 and Fyn**  
Qi Shi, Chinese Center for Disease Control and Prevention
- 29. Infection of prions and treatment of PrP106-126 downregulate the endogenous levels of protein 14-3-3 and trigger the mitochondrial apoptosis possibly via activating Bax pathway**  
Qi Shi, Chinese Center for Disease Control and Prevention
- 30. Abundant intracellular and neuritic autophagic vacuoles in hippocampus of PrP-deficient mice**  
Hae-Young Shin, Hallym University
- 31. TSE amplification between wild type mouse by *in vitro* conversion of prion protein**  
Hyun-Joo Sohn, Animal and Plant Quarantine Agency
- 32. The mechanism of asymmetric distribution of prion proteins in the yeast cell**  
Genjiro Suzuki, RIKEN
- 33. An autopsy case of familial Creutzfeldt-Jakob disease with M232R-129M/V**  
Mikiko Tada, Yokohama City University
- 34. Mechanism underlying difference between transmissible prions and nontransmissible amyloids**  
Hideki Taguchi, Tokyo Institute of Technology
- 35. A family and patient guidebook about genetic testing of prion disease**  
Chieko Tamura, Kiba Park Clinic / Juntendo University Hospital
- 36. Structural basis for conformational plasticity of yeast prion amyloid**  
Motomasa Tanaka, RIKEN
- 37. Comparative analysis of gene expression profiles between cortex and thalamus in Chinese fatal familial insomnia patients**  
Chan Tian, Chinese Center for Disease Control and Prevention
- 38. SIRT1 activity is associated with cellular PrP<sup>Sc</sup> level in scrapie agent 263K-infected hamster and SMB.s15 cells**  
Jing Wang, Chinese Center for Disease Control and Prevention

**39. Nearly reversible conformational change of H2 amyloid fibrils as revealed by pH-jump experiments**

Kei-ichi Yamaguchi, Gifu University

**40. A cation exchanger specifically enhances cell-protein misfolding cyclic amplification of variant Creutzfeldt-Jakob disease prion despite its weaker adsorbing ability**

Takashi Yokoyama, Japan Blood Products Organization

**41. Defining Korean Creutzfeldt-Jakob disease strains and their transmissible properties**

Yun-Jung Lee, Hallym University

**42. Quantitative analysis of seeding activity in human prion disease materials using end-point RT-QUIC**

Hanae Takatsuki, Nagasaki University

**43. Transient PrP<sup>Sc</sup> propagation in dead cells**

Kenichi Takahashi, Gifu University

**44. The role of Aldolase A in transcytosis of PrP<sup>Sc</sup> by intestinal M cells**

Yuya Nagasawa, Tohoku University

**45. Analysis of Interferon Regulatory Factor-3 (IRF-3) Promoter: Implications for Prion Infection**

Takujiro Homma, Nagasaki University

**46. p62/SQSTM1 promotes the sequestration and clearance of pathogenic prion protein**

Takujiro Homma, Nagasaki University

**47. New method for discovery of novel anti-prion compounds: Intermediate structure-based drug design (IBDD)**

Biao Ma, Gifu University

**48. Possible involvement of a novel mitochondrial quality control system for the PrP<sup>C</sup>-dependent neuronal cell death**

Kana Miyashita, Tokyo Medical University

**49. Experimental transmission of AA amyloidosis in vaccinated white hens**

Tomoaki Murakami, Gifu University

**50. The effect of metal ions on the structural stability of prion protein**

Kaori Ubagai, Nagasaki University