



CURRICULUM VITAE

Personal Information

Name: Huifang Shang
Sex: Female
Date of Birth: June 7, 1970
Place of Birth: Shanxi, China
Nationality: Chinese
Health: Excellent
Address: Department of Neurology, West China Hospital, Sichuan University, Chengdu, Sichuan, China
Phone: +86-28-85422548(office); +86-28-66310568 (home)
Email: hfshang@yahoo.com

Education

2002 - 2003 **Medical Doctor**, University of Bern, Bern, Switzerland, 2003
MD thesis: *Molecular and genetic analysis in primary dystonia*

1996 - 1999 **Master of Medical Science**, West China University of Medical Sciences, Chengdu, China, 1999
Dissertation: *Protective and superimposing effects of Estradiol, Breviscapini, Lumbrokinase on focal cerebral ischemic-reperfusion injury in rats*

1988 - 1993 **Bachelor of Medical Science**, Suzhou Medical College, Suzhou, China, 1993

Professional Experience

2011.7- **Professor** in the Dept. of Neurology, West China, Hospital, Sichuan, University, Chengdu, China

- 2009- 2011 **Associate professor** in the Dept. of Neurology, West China, Hospital, Sichuan, University, Chengdu, China
- 2008-2009 **Research fellow** in the Dept. of Pharmacology, Texas University, SanAntonio, Texas, United States
- 2006 - 2008 **Associate professor** in the Dept. of Neurology, West China Hospital, Sichuan University, Chengdu, China
In charge of teaching medical students, residents, trained doctors in neurology. Supervise graduate students
- 2004 - 2006 **Attending physician** in the Dept. of Neurology, West China Hospital, Sichuan University, Chengdu, China
In charge of teaching medical students, residents, trained doctors neurology
- 2002 - 2004 **Research Fellow** in the Neuromorphologic Laboratory, Dept. of Neurology, Inselspital, Bern University, Bern, Switzerland
Sponsored by IBRO; Project: Molecular genetic investigation of dystonia in a Swiss and Chinese population
- 1997 - 2001 **Resident, chief resident and attending physician** in the Dept. of Neurology, West China Hospital, Chengdu, China
In charge of teaching interns neurology
Research project sponsored by Chinese Ministry of Health; Project: Multicentric Randomised Controlled Trial of Patients with Acute Cerebral Infarction Treated with Cyclophosphamide and Colchicine
- 1993 - 1996 **Resident** in Dept. of Neurology, Chengdu No. 416 Hospital, Chengdu, Sichuan, China

Publications List

Refereed Journal Papers in English

- 1) Chen Y, Wei QQ, Ou R, Cao B, Chen X, Zhao B, Guo X, Yang Y, Chen K, Wu Y, Song W, Shang HF. Genetic Variants of SNCA Are Associated with Susceptibility to Parkinson's Disease but Not Amyotrophic Lateral Sclerosis or Multiple System Atrophy in a Chinese Population. PLoS One. 2015 Jul 24;10 (7):e0133776.
(Corresponding author)

- 2) Chen X, Chen Y, Wei Q, Guo X, Cao B, Ou R, Zhao B, Shang HF. Assessment of TREM2 rs75932628 association with amyotrophic lateral sclerosis in a Chinese population. *J Neurol Sci*. 2015 Aug 15;355(1-2):193-5. **(Corresponding author)**
- 3) Luo C, Guo X, Song W, Chen Q, Yang J, Gong Q, Shang HF. The trajectory of disturbed resting-state cerebral function in Parkinson's disease at different Hoehn and Yahr stages. *Hum Brain Mapp*. 2015 Aug;36(8):3104-16. **(Corresponding author)**
- 4) Luo CY, Guo XY, Song W, Chen Q, Cao B, Yang J, Gong QY, Shang HF. Functional connectome assessed using graph theory in drug-naive Parkinson's disease. *J Neurol*. 2015 Jun;262(6):1557-67. **(Corresponding author)**
- 5) Ou R, Guo X, Wei Q, Cao B, Yang J, Song W, Chen K, Zhao B, Chen X, Shang H. Diurnal drooling in Chinese patients with Parkinson's disease. *J Neurol Sci*. 2015 Jun 15;353(1-2):74-8. **(Corresponding author)**
- 6) Ou R, Guo X, Wei Q, Cao B, Yang J, Song W, Shao N, Zhao B, Chen X, Shang H. Prevalence and clinical correlates of drooling in Parkinson disease: a study on 518 Chinese patients. *Parkinsonism Relat Disord*. 2015 Mar;21(3):211-5. **(Corresponding author)**
- 7) Guo X, Song W, Chen K, Chen X, Zheng Z, Cao B, Huang R, Zhao B, Wu Y, Shang HF. Associations between neuropsychiatric symptoms and cognition in Chinese idiopathic Parkinson's disease patients. *J Clin Neurosci*. 2015 Mar;22(3):578-82. **(Corresponding author)**
- 8) Wei Q, Chen X, Zheng Z, Guo X, Huang R, Cao B, Zeng Y, Shang H. The predictors of survival in Chinese amyotrophic lateral sclerosis patients. *Amyotroph Lateral Scler Frontotemporal Degener*. 2015 Jun;16(3-4):237-44. **(Corresponding author)**
- 9) Yuan X, Chen Y, Cao B, Zhao B, Wei Q, Guo X, Yang Y, Yuan L, Shang H. An association analysis of the R1628P and G2385R polymorphisms of the LRRK2 gene in multiple system atrophy in a Chinese population. *Parkinsonism Relat Disord*. 2015 Feb;21(2):147-9. **(Corresponding author)**
- 10) Shao N, Yang J, Shang H. Voxelwise meta-analysis of gray matter anomalies in Parkinson variant of multiple system atrophy and Parkinson's disease using anatomic likelihood estimation. *Neurosci Lett*. 2015 Feb 5;587:79-86. **(Corresponding author)**

- 11) Chen YP, Zhao B, Cao B, Song W, Guo X, Wei QQ, Yang Y, Yuan LX, Shang HF. Mutation scanning of the COQ2 gene in ethnic Chinese patients with multiple-system atrophy. *Neurobiol Aging*. 2015 Feb;36(2):1222.e7-11. (**Corresponding author**)
- 12) Wei Q, Chen X, Zheng Z, Huang R, Guo X, Cao B, Bak TH, Shang H. Screening for cognitive impairment in a Chinese ALS population. *Amyotroph Lateral Scler Frontotemporal Degener*. 2015 Mar;16(1-2):40-5. (**Corresponding author**)
- 13) Chen X, Feng W, Huang R, Guo X, Chen Y, Zheng Z, Shang H. Evidence for peripheral immune activation in amyotrophic lateral sclerosis. *J Neurol Sci*. 2014 Dec 15;347(1-2):90-5. (**Corresponding author**)
- 14) Chen D, Guo X, Zheng Z, Wei Q, Song W, Cao B, Huang R, Yang R, Huifang S. Depression and anxiety in ALS: Correlations between the distress of patients and caregivers. *Muscle Nerve*. 2015 Mar; 51(3):353-7. (Co-**Corresponding author**)
- 15) Wei Q, Chen X, Zheng Z, Huang R, Guo X, Cao B, Zhao B, Shang HF. Frontal lobe function and behavioral changes in amyotrophic lateral sclerosis: a study from Southwest **China**. *J Neurol*. 2014 Dec; 261(12): 2393-400 (**Corresponding author**)
- 16) Guo XY, Chen YP, Song W, Zhao B, Cao B, Wei QQ, Ou RW, Yang Y, Yuan LX, Shang HF. SNCA variants rs2736990 and rs356220 as risk factors for Parkinson's disease but not for amyotrophic lateral sclerosis and multiple system atrophy in a Chinese population. *Neurobiol Aging*. 2014 Dec; 35(12):2882.e1-6 (**Corresponding author**)
- 17) Ou R, Guo X, Song W, Cao B, Yang J, Wei Q, Shao N, Shang H. Freezing of gait in Chinese patients with Parkinson Disease. *J Neurol Sci*. 2014 Oct 15;345(1-2):56-60. (**Corresponding author**)
- 18) Guo XY, Chen YP, Song W, Zhao B, Cao B, Wei QQ, Ou RW, Yang Y, Yuan LX, Shang HF. An association analysis of the rs1572931 polymorphism of the RAB7L1 gene in Parkinson's disease, amyotrophic lateral sclerosis and multiple system atrophy in **China**. *Eur J Neurol*. 2014 Oct; 21(10):1337-43. (**Corresponding author**)
- 19) Zhao B, Wei Q, Wang Y, Chen Y, Shang H. Posterior reversible encephalopathy syndrome in acute intermittent porphyria. *Pediatr Neurol*. 2014 Sep; 51(3):457-60.

- 20) Ou R, Guo X, Song W, Cao B, Wei Q, Shao N, Zhao B, **Shang H**. Characteristics of non-motor symptoms in patients with Parkinson's disease exhibiting camptocormia. *Gait Posture*. 2014 Jul;40(3):447-50. **(Corresponding author)**
- 21) Chen X, Guo X, Huang R, Chen Y, Zheng Z, **Shang H**. Serum uric acid levels in patients with Alzheimer's disease: a meta-analysis. *PLoS One*. 2014 Apr 8;9(4):e94084. **(Corresponding author)**
- 22) Wu Y, Guo XY, Wei QQ, Song W, Chen K, Cao B, Ou RW, Zhao B, **Shang HF**. Determinants of the quality of life in Parkinson's disease: results of a cohort study from Southwest **China**. *J Neurol Sci*. 2014 May 15;340(1-2):144-9. **(Corresponding author)**
- 23) Shao N, Yang J, Li J, **Shang HF**. Voxelwise meta-analysis of gray matter anomalies in progressive supranuclear palsy and Parkinson's disease using anatomic likelihood estimation. *Front Hum Neurosci*. 2014 Feb 18;8:63. **(Corresponding author)**
- 24) Cao B, Guo X, Chen K, Song W, Huang R, Wei QQ, Zhao B, **Shang HF**. Serum lipid levels are associated with the prevalence but not with the disease progression of multiple system atrophy in a Chinese population. *Neurol Res*. 2014 Feb;36(2):150-6.
- 25) Zheng Z, Guo X, Wei Q, Song W, Cao B, Huang R, Ou R, Chen X, **Shang H**. Serum uric acid level is associated with the prevalence but not with survival of amyotrophic lateral sclerosis in a Chinese population. *Metab Brain Dis*. 2014 Sep;29(3):771-5. **(Corresponding author)**
- 26) Wei QQ, Chen Y, Zheng ZZ, Chen X, Huang R, Yang Y, Burgunder J, **Shang HF**. Spastin mutation screening in Chinese patients with pure hereditary spastic paraplegia. *Parkinsonism Relat Disord*. 2014 Aug;20(8):845-9. **(Corresponding author)**
- 27) Yang J, Luo C, Song W, Guo X, Zhao B, Chen X, Huang X, Gong Q, **Shang HF**. Diffusion tensor imaging in blepharospasm and blepharospasm-oromandibular dystonia. *J Neurol*. 2014 Jul;261(7):1413-24. **(Co-Corresponding author)**
- 28) Chen X, Guo X, Huang R, Zheng Z, Chen Y, **Shang HF**. An exploratory study of serum creatinine levels in patients with amyotrophic lateral sclerosis. *Neurol Sci*. 2014 Oct;35(10):1591-7. **(Corresponding author)**

- 29) Li J, Luo C, Chen Y, Chen Q, Huang R, Sun J, Gong Q, Wu X, Qi Z, Liang Z, Li L, Li H, Li P, Wang W, **Shang HF**. Parkinson's disease-related modulation of functional connectivity associated with the striatum in the resting state in a nonhuman primate model. *Brain Res.* 2014 Mar 25;1555:10-9. (**Corresponding author**)
- 30) Li J, Chen D, Song W, Chen K, Cao B, Huang R, Yang R, **Shang H**. Survey on general knowledge on Parkinson's disease in patients with Parkinson's disease and current clinical practice for Parkinson's disease among general neurologists from Southwest China. *Clin Neurol Neurosurg.* 2014 Mar;118:16-20. (**Co-Corresponding author**)
- 31) Zheng Z, Guo X, Huang R, Chen X, **Shang H**. An exploratory study of the association between thyroid hormone and survival of amyotrophic lateral sclerosis. *Neurol Sci.* 2014 Jul;35(7):1103-8. (**Corresponding author**)
- 32) Chen X, Huang R, Chen Y, Zheng Z, Chen K, Song W, Zhao B, Yang Y, Yuan L, **Shang H**. Association analysis of four candidate genetic variants with sporadic amyotrophic lateral sclerosis in a Chinese population. *Neurol Sci.* 2014 Jul;35(7):1089-95. (**Corresponding author**)
- 33) Song W, Guo X, Chen K, Huang R, Zhao B, Cao B, Chen Y, **Shang HF**. Camptocormia in Chinese patients with Parkinson's disease. *J Neurol Sci.* 2014 Feb 15; 337(1-2):173-5. (**Corresponding author**)
- 34) Luo C, Chen Q, Song W, Chen K, Guo X, Yang J, Huang X, Gong Q, **Shang HF**. Resting-state fMRI study on drug-naive patients with Parkinson's disease and with depression. *J Neurol Neurosurg Psychiatry.* 2014, 85(6):675-83. (**Co-Corresponding author**)
- 35) Song W, Guo X, Chen K, Chen X, Cao B, Wei Q, Huang R, Zhao B, Wu Y, **Shang HF**. The impact of non-motor symptoms on the Health-Related Quality of Life of Parkinson's disease patients from Southwest China. *Parkinsonism Relat Disord.* 2014, 20 (2): 149-52. (**Corresponding author**)
- 36) Chen Y, Zheng ZZ, Chen X, Huang R, Yang Y, Yuan L, Pan L, Hadano S, **Shang HF**. SQSTM1 mutations in Han Chinese populations with sporadic amyotrophic lateral sclerosis. *Neurobiol Aging.* 2014, 35(3): 726.e7-9. (**Corresponding author**)

- 37) Yang J, Shao N, Li J, **Shang H**. Voxelwise meta-analysis of white matter abnormalities in progressive supranuclear palsy. *Neurol Sci*. 2014, 35(1): 7-14. **(Corresponding author)**
- 38) Luo C, Song W, Chen Q, Zheng Z, Chen K, Cao B, Yang J, Li J, Huang X, Gong Q, **Shang HF**. Reduced functional connectivity in early-stage drug-naive Parkinson's disease: a resting-state fMRI study. *Neurobiol Aging*. 2014, 35(2):431-41. **(Co-Corresponding author)**
- 39) Cao B, Tang Y, Li J, Zhang X, **Shang HF**, Zhou D. A meta-analysis of voxel-based morphometry studies on gray matter volume alteration in juvenile myoclonic epilepsy. *Epilepsy Res*. 2013 Oct;106(3):370-7. **(Corresponding author)**
- 40) Yang J, Luo C, Song W, Chen Q, Chen K, Chen X, Huang X, Gong Q, **Shang H**. Altered regional spontaneous neuronal activity in blepharospasm: a resting state fMRI study. *J Neurol*. 2013 Nov;260(11):2754-60. **(Corresponding author)**
- 41) Guo X, Song W, Chen K, Chen X, Zheng Z, Cao B, Huang R, Zhao B, Wu Y, **Shang HF**. Gender and onset age-related features of non-motor symptoms of patients with Parkinson's disease - A study from Southwest **China**. *Parkinsonism Relat Disord*. 2013 Nov;19(11):961-5. **(Corresponding author)**
- 42) Chen Y, Song W, Yang J, Chen K, Huang R, Zhao B, Cao B, Burgunder J, **Shang HF**. Association of the Val66Met polymorphism of the BDNF gene with primary cranial-cervical dystonia patients from South-west China. *Parkinsonism Relat Disord*. 2013 Nov;19(11):1043-5. **(Corresponding author)**
- 43) Cao B, Guo X, Chen K, Song W, Huang R, Wei QQ, Zhao B, **Shang HF**. Uric acid is associated with the prevalence but not disease progression of multiple system atrophy in Chinese population. *J Neurol*. 2013 Oct;260(10):2511-5. **(Corresponding author)**
- 44) Chen YP, Song W, Huang R, Chen K, Zhao B, Li J, Yang Y, Shang HF. GAK rs1564282 and DGKQ rs11248060 increase the risk for Parkinson's disease in a Chinese population. *J Clin Neurosci*. 2013 Jun;20(6):880-3. **(Corresponding author)**
- 45) Guo X, Song W, Chen K, Chen X, Zheng Z, Cao B, Huang R, Zhao B, Wu Y, Shang HF. Disease duration-related differences in non-motor symptoms: A study of 616

Chinese Parkinson's disease patients. *J Neurol Sci.* 2013 Jul 15;330(1-2):32-7.
(**Corresponding author**)

- 46) Guo XY, Cao B, Lei F, Huang L, Chen K, Song W, Zhao B, Tang X, Shang H. Clinical and polysomnographic features of patients with multiple system atrophy in Southwest China. *Sleep Breath.* 2013 Dec;17(4):1301-7. (**Corresponding author**)
- 47) Luo C, Chen Y, Song W, Chen Q, Gong Q, Shang HF. Altered intrinsic brain activity in patients with paroxysmal kinesigenic dyskinesia by PRRT2 mutation: Altered brain activity by PRRT2 mutation. *Neurol Sci.* 2013 Nov;34(11):1925-31.
(**Corresponding author**)
- 48) Chen YP, Song W, Yang J, Zheng ZZ, Huang R, Chen K, Zhao B, Chen XP, Burgunder JM, Shang HF. PRRT2 mutation screening in patients with paroxysmal kinesigenic dyskinesia from Southwest China. *Eur J Neurol.* 2014, 21(1): 174-6.
(Corresponding author)
- 49) Chen Y, Zheng ZZ, Huang R, Chen K, Song W, Zhao B, Chen X, Yang Y, Yuan L, Shang HF. PFN1 mutations are rare in Han Chinese populations with amyotrophic lateral sclerosis. *Neurobiol Aging.* 2013 Jul;34(7):1922.e1-5. (Corresponding author)
- 50) Chen Y, Chen K, Song W, Chen X, Cao B, Huang R, Zhao B, Guo X, Burgunder J, Li J, Shang HF. VPS35 Asp620Asn and EIF4G1 Arg1205His mutations are rare in Parkinson disease from Southwest **China**. *Neurobiol Aging.* 2013 Jun;34(6):1709.e7-8. (**Corresponding author**)
- 51) Zheng Z, Sheng L, **Shang H**. Statins and amyotrophic lateral sclerosis: a systematic review and meta-analysis. *Amyotroph Lateral Scler Frontotemporal Degener.* 2013 May;14(4):241-5 (**Corresponding author**)
- 52) Chen Y, Chen K, Burgunder JM, Song W, Huang R, Zhao B, Cao B, Chen X, Jiang Y, **Shang HF**. Association of rs1182 polymorphism of the DYT1 gene with primary dystonia in Chinese population. *J Neurol Sci.* 2012 Dec 15;323(1-2):228-31.
(**Corresponding author**)
- 53) Luo C, Chen Q, Huang R, Chen X, Chen K, Huang X, Tang H, Gong Q, **Shang HF**. Patterns of spontaneous brain activity in amyotrophic lateral sclerosis: a resting-state FMRI study. *PLoS One.* 2012;7(9):e45470. (**Corresponding author**)

- 54) Chen Y, Huang R, Chen K, Song W, Yang Y, Zhao B, Li J, **Shang HF**. Association analysis of PON polymorphisms in sporadic ALS in a Chinese population. *Neurobiol Aging*. 2012 Dec;33(12):2949.e1-3. **(Corresponding author)**
- 55) Chen Y, Zeng Y, Huang R, Yang Y, Chen K, Song W, Zhao B, Li J, Yuan L, **Shang HF**. No association of five candidate genetic variants with amyotrophic lateral sclerosis in a Chinese population. *Neurobiol Aging*. 2012 Nov;33(11):2721.e3-5. **(Corresponding author)**
- 56) Zhao B, Song W, Chen YP, Huang R, Chen K, Cao B, Yang Y, **Shang HF**. Association Analysis of Single-Nucleotide Polymorphisms of USP24 and USP40 with Parkinson's Disease in the Han Chinese Population. *Eur Neurol*. 2012;68(3):181-4. **(Corresponding author)**
- 57) Yang J, Pan P, Song W, **Shang HF**. Quantitative meta-analysis of gray matter abnormalities in semantic dementia. *J Alzheimers Dis*. 2012 Jan 1;31(4):827-33. **(Corresponding author)**
- 58) Song W, Chen YP, Huang R, Chen K, Pan PL, Li J, Yang Y, **Shang HF**. GLIS1 rs797906: An Increased Risk Factor for Late-Onset Parkinson's Disease in the Han Chinese Population. *Eur Neurol*. 2012;68(2):89-92. **(Corresponding author)**
- 59) Chen K, Chen YP, Song W, Huang R, Zhao B, Cao B, Yang Y, Satake W, Toda T, **Shang HF**. Association analysis of LRP8 SNP rs3820198 and rs5174 with Parkinson's disease in Han Chinese population. *Neurol Res*. 2012 Sep;34(7):725-9. **(Corresponding author)**
- 60) Pan PL, Song W, Yang J, Huang R, Chen K, Gong QY, Zhong JG, Shi HC, **Shang HF**. Gray matter atrophy in behavioral variant frontotemporal dementia: a meta-analysis of voxel-based morphometry studies. *Dement Geriatr Cogn Disord*. 2012;33(2-3):141-8. **(Corresponding author)**
- 61) Li J, Pan P, Song W, Huang R, Chen K, **Shang H**. A meta-analysis of diffusion tensor imaging studies in amyotrophic lateral sclerosis. *Neurobiol Aging*. 2012, 33(8):1833-8. **(Corresponding author)**
- 62) Chen Y, Burgunder JM, Song W, Huang R, **Shang HF**. Assessment of D216H DYT1 polymorphism in a Chinese primary dystonia patient cohort. *Eur J Neurol*. 2012, 19(6):924-6 **(Corresponding author)**

- 63) Li P, Huang R, Song W, Ji J, Burgunder JM, Wang X, Zhong Q, Kaelin-Lang A, Wang W, **Shang HF**. Deep brain stimulation of the globus pallidus internal improves symptoms of chorea-acanthocytosis. *Neurol Sci.* 2012, 33(2):269-74. **(Corresponding author)**
- 64) Huang R, Fang DF, Ma MY, Guo XY, Zhao B, Zeng Y, Zhou D, Yang Y, **Shang HF**. TARDBP gene mutations among Chinese patients with sporadic amyotrophic lateral sclerosis. *Neurobiol Aging.* 2012, 33(5):1015.e1-6. **(Corresponding author)**
- 65) Yang J, Pan P, Song W, Huang R, Li J, Chen K, Gong Q, Zhong J, Shi H, **Shang H**. Voxelwise meta-analysis of gray matter anomalies in Alzheimer's disease and mild cognitive impairment using anatomic likelihood estimation. *J Neurol Sci.* 2012, 316, 21-29. **(Corresponding author)**
- 66) Zheng Z, Pan P, Wang W, **Shang H**. Neural network of primary focal dystonia by an anatomic likelihood estimation meta-analysis of gray matter abnormalities. *J Neurol Sci.* 2012, 316, 51-55. **(Corresponding author)**
- 67) Wei Song, YongPing Chen, Rui Huang, Ke Chen, Ping Lei Pan, Jianpeng Li, Yuan Yang1, **Hui-Fang Shang**. Analysis of the genotype of diacylglycerol kinase delta single-nucleotide polymorphisms in Parkinson disease in the Han Chinese population. *Neurology India.* 2012, 60 (1): 82—5. **(Corresponding author)**
- 68) Li J, Pan P, Huang R, **Shang H**. A meta-analysis of voxel-based morphometry studies of white matter volume alterations in Alzheimer's disease. *Neurosci Biobehav Rev.* 2012, 36(2):757-63. **(Corresponding author)**
- 69) **Shang H**, Danek A, Landwehrmeyer B, Burgunder JM. Huntington's disease: new aspects on phenotype and genotype. *Parkinsonism Relat Disord.* 2012, 18 Suppl 1:S107-9. **(Co-author)**
- 70) Li J, Zhang Z, **Shang H**. A meta-analysis of voxel-based morphometry studies on unilateral refractory temporal lobe epilepsy. *Epilepsy Res.* 2012, 98: 97—103. **(Corresponding author)**
- 71) Pan PL, Song W, **Shang HF**. Voxel-wise meta-analysis of gray matter abnormalities in idiopathic Parkinson's disease. *Eur J Neurol.* 2012, 19(2):199-206. **(Corresponding author)**
- 72) Chen Y, Huang R, Yang Y, Chen K, Song W, Pan P, Li J, **Shang HF**. Ataxin-2 intermediate-length polyglutamine: a possible risk factor for Chinese patients with

amyotrophic lateral sclerosis. *Neurobiol Aging*. 2011, 32(10):1925.e1-5.
(**Corresponding author**)

- 73) Song W, Chen Y, Huang R, Chen K, Pan P, Yang Y, **Shang HF**. Novel THAP1 gene mutations in patients with primary dystonia from Southwest **China**. *J Neurol Sci*. 2011, 309(1-2):63-7. (**Corresponding author**)
- 74) Pan PL, Tang HH, Chen Q, Song W, **Shang HF**. Desferrioxamine treatment of aceruloplasminemia: Long-term follow-up. *Mov Disord*. 2011, 26(11):2142-4. (**Corresponding author**)
- 75) Yang R, Huang R, Chen D, Song W, Zeng Y, Zhao B, Zhou D, **Shang HF**. Causes and places of death of patients with amyotrophic lateral sclerosis in south-west China. *Amyotroph Lateral Scler*. 2011, 12(3):206-9. (**Corresponding author**)
- 76) Li P, Mao B, **Shang H**, Kaelin-Lang A, Wang W. Pilot study on a fast postoperative programming approach to subthalamic nucleus stimulation in Parkinson's disease. *Neurol India*. 2011, 59(5):669-73. ([Co-author](#))
- 77) He X, Lu X, Hu J, Xi J, Zhou D, **Shang H**, Liu L, Zhou H, Yan B, Yu L, Hu F, Liu Z, He L, Yao X, Xu Y. H63D polymorphism in the hemochromatosis gene is associated with sporadic amyotrophic lateral sclerosis in China. *Eur J Neurol*. 2011, 18(2):359-61. ([Co-author](#))
- 78) Liu X, Zhang SS, Fang DF, Ma MY, Guo XY, Yang Y, **Shang HF**. GCH1 Mutation and Clinical Study of Chinese Patients with Dopa-responsive Dystonia. *Movement disorder*. 2010, 25(4):447-454. (**Corresponding author**)
- 79) Zhang SS, Fang DF, Burgunder JM, Chen XP, Zhang YW, **Shang HF**. Clinical feature and DYT1 mutation screening in primary dystonia patients from South-West China. *European J Neurology*. 2010, 17: 846-851. (**Corresponding author**)
- 80) Zhou B, Chen Q, Zhang Q, Chen L, Gong Q, **Shang H**, Tang H, Zhou D. Hyperactive putamen in patients with paroxysmal kinesigenic choreoathetosis: a resting-state functional magnetic resonance imaging study. *Mov Disord*. 2010, 25(9):1226-31. ([Co-author](#))
- 81) Yang TH, Tian LY, **Shang HF**, Cheng XW, Geng J, Chen L, Zhou D. Suppression of the multidrug transporter P-glycoprotein using RNA interference in cultured rat astrocytes induced by coriaria lactone. *Neurol Res*. 2009, 31(10):1084-91. ([Co-author](#))

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- 83) Hu XH, Chen XP, Huang R, **Shang Huifang**. Role of DYT1 gene in early-onset primary torsin dystonia. *Neural Regen Res*, 2010, 5(18):1429-1435. **(corresponding author)**
- 84) Guo XY, Zhang SS, Burgunder JM, **Shang HF**. Clinical features of Huntington disease in 243 Chinese patients. *Neural Regen Res*. 2010, 5(2):102-107. **(Corresponding author)**
- 85) Fang DF, Zhang SS, Guo XY, Zeng Y, Yang Y, Zhou D, **Shang HF**. Clinical and genetic features of patients with sporadic amyotrophic lateral sclerosis in south-west China. *Amyotroph Lateral Scler*. 2009, 10(1):1-6. **(Co-Corresponding author)**.
- 86) XP Chen, **HF Shang**, ZM Luo. Genetic classification and molecular mechanisms of primary dystonia. *Neural Regen Res*, 2008, 3(3), 296-300. **(Corresponding author)**.
- 87) YW Zhang, SS Zhang, **HF Shang**. Clinical characteristics of myotonia congenita in China: Literature analysis of the past 30 years. *Neural Regen Res*, 2008, 3(2), 216-20. **(Corresponding author)**.
- 88) Li Q, **Shang H**, Zhou D, Liu R, He L, Zheng H. Repeated embolism and multiple aneurysms: central nervous system manifestations of cardiac myxoma. *Eur J Neurol*. 2008 Dec;15(12):e112-3 **(Co-author)**
- 89) SS Wu, **HF Shang**, XY Zou. Dopaminergic system abnormalities: Etiopathogenesis of dystonia. *Neural Regen Res*, 2008, 3(3), 301-4. **(Co-author)**
- 90) Q Chen, ZP Yao, D Zhou, HB Zheng, **HF Shang**. Lateral sinus thrombosis and intracranial hypertension associated with primary hypothyroidism. *Neuro Endocrinol Lett*. 2008; 29(1):41-3. **(Corresponding author)**.
- 91) Q Chen, S Lui, JG Wang, L Ou-Yang, D Zhou, JM Burgunder, QY Gong, **HF Shang**. Diffusion tensor imaging of two unrelated Chinese men with hereditary spastic paraplegia associated with thin corpus callosum. *Neuroscience Letters*, 2008, 441(1): 21-24. **(corresponding author)**
- 92) XP Chen, YW Zhang, SS Zhang, Q Chen, JM Burgunder, SH Wu, Y Yang, ZM Luo,

HF Shang. A novel mutation of epsilon-sarcoglycan gene in a Chinese family with myoclonus-dystonia syndrome. *Movement Disorders*, 2008,23(10): 1472-1475.

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- 93) Q Chen, XP Chen, L Zou, D Zhou, QY Gong, JM Burgunder, **HF Shang**. High brain iron level in asymptomatic carriers of heterozygous ceruloplasmin gene mutations. *Movement Disorders*, 2008, 23(6):916-917. (Corresponding author)
- 94) SS Zhang, Q Chen, XP Chen, JG Wang, JM Burgunder, Y Yang, **HF Shang**. Two novel mutations in the *SPG11* gene causing hereditary spastic paraplegia associated with thin corpus callosum. *Movement Disorders*, 2008,23(6):918-919.(Corresponding author)
- 95) McNeill A, Pandolfo M, Kuhn J, **Shang H**, Miyajima H. The neurological presentation of ceruloplasmin gene mutations. *European Neurology*, 2008, 60(4):200-205. (Co-author)
- 96) Chen Q, Lui S, Li CX, Jiang LJ, Ou-Yang L, Tang HH, **Shang HF**, Huang XQ, Gong QY, Zhou D. MRI-negative refractory partial epilepsy: Role for diffusion tensor imaging in high field MRI. *Epilepsy Research*, 2008, 80(1):83-89. (Co-author)
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Grants

2005 - 2007 Principal investigator

Project: *The role of vesicle membrane proteins, cytoskeleton proteins and ion channels in the mechanism of DYT1 and SGCE gene involved in dystonia by using small RNA interference inhibited the expression of DYT1 and SGCE gene in neurons.*

Funded by: *National Natural Science Foundation of China*

2010 - 2012 Principal investigator

Project: *Study on abnormal function and structural neuronal network in MPTP-induced chronic PD rhesus model*

Funded by: *National Natural Science Foundation of China*

2014 - 2017 Principal investigator

Project: *MiRNA in the pathogenesis of amyotrophic lateral sclerosis*

Funded by: *National Natural Science Foundation of China*

2016 - 2019 Principal investigator

Project: *Imaging in genetics in in cognitive impairment of Parkinson ' s disease*

Funded by: *National Natural Science Foundation of China*