

RESEARCH

Open Access

Prevalence of wearing-off and dyskinesia among the patients with Parkinson's disease on levodopa therapy: a multi-center registry survey in mainland China

Wei Chen^{1†}, Qin Xiao^{1†}, Ming Shao², Tao Feng³, Wei-Guo Liu⁴, Xiao-Guang Luo⁵, Xiao-Chun Chen⁶, An-Mu Xie⁷, Chun-Feng Liu⁸, Zhen-Guo Liu⁹, Yi-Ming Liu¹⁰, Jian Wang¹¹ and Sheng-Di Chen^{1*}

Abstract

Objective: Chronic levodopa (L-dopa) treatment in Parkinson's disease (PD) is often associated with the development of motor complications, but the corresponding epidemiological data is rare in Chinese PD patients. The present survey was to investigate the prevalence rate of wearing-off (WO) and dyskinesia among the patients with PD in China.

Methods: From May 2012 to October 2012, a 3-step registry survey for wearing off (WO) and dyskinesia patients with PD receiving levodopa therapy was performed simultaneously at 28 movement disorders clinics in China.

Results: There were 1,558 PD patients fulfilling the inclusion criteria. Among them, 1,051 had at least one positive response of 9-item wearing off questionnaire (WOQ-9), 724 and 160 patients were finally diagnosed with WO and dyskinesia by movement disorders specialists, respectively. The overall prevalence rates of WO and dyskinesia were 46.5% (95% CI 44.0% - 48.9%) and 10.3% (95% CI 8.8% - 11.8%), respectively. The mean score of WOQ-9 for those with WO was 3.8 (SD = 1.8), with movement slowness being the most common motor symptoms and pain/aching being the most common non-motor symptoms. Better improvement of motor symptoms ($n = 354$, 87.8%) and long-term disease control and drug selection ($n = 288$, 71.5%) were the two most frequently considered factors when movement disorders specialists adjusted therapeutic strategies for patients with WO.

Conclusions: This survey provided the first multi-center epidemiological data of motor complications among PD patients on L-dopa therapy from mainland China. WO prevalence rate among Chinese PD patients was in line with, while dyskinesia prevalence rate was lower than previous reports from other Countries.

Keywords: Parkinson's disease, Wearing-off, Dyskinesia, Epidemiology

Introduction

Up to now, Levodopa (L-dopa) is still recognized as the most widely used and effective medication for Parkinson's disease (PD), but long period therapy is often associated with the development of motor complications such as wearing-off (WO) and dyskinesia, which brought not only challenges for neurologists, but also impaired daily living and poor life quality of PD patients. Generally, the first

motor complication is predictable WO in advancing PD, that is, a recurrence of motor and non-motor symptoms preceding scheduled doses of anti-PD medication that usually improved after the next dosage [1]. Early identification of such condition is of great importance for the timely optimized treatment of PD.

Cumulative evidence indicated that approximately 40% patients treated with L-dopa for 4–6 years experienced motor complications in Western Countries [2,3], whereas, the corresponding epidemiological data was rare in Chinese PD patients. Heterogeneity exists in previous reports from Hong Kong [4] and Suzhou [5], China. Moreover, both of them were based on single center data with limited sample

* Correspondence: chen_sd@medmail.com.cn

†Equal contributors

¹Department of Neurology, Rui Jin Hospital affiliated to Shanghai Jiao Tong University School of Medicine, Rui Jin 2nd Road 197, Shanghai 200025, China
Full list of author information is available at the end of the article

size. Thus, the true nature of motor complications, especially WO in Chinese PD patients deserves further investigation. Since the 9-item wearing-off questionnaire (WOQ-9) is a simple and highly-sensitive diagnostic screening scale for WO [6,7], we considered that WO was rarely present in patients with zero point on WOQ-9 as recommended by Movement Disorders Society (MDS) in 2011 [8].

Once recognized, WO can be effectively managed via a number of therapeutic options, including adjustments of L-dopa dosage and dosage form, adjunctive therapy with catechol-O-methyltransferase (COMT) inhibitors, monoamine oxidase-B (MAO-B) inhibitors and dopamine agonists. Nevertheless, clinical experience and treatment concepts from movement disorders specialists also play critical roles in the optimized therapy of WO [9,10]. Since PD is still an incurable disease, current therapy could only improve the symptoms, the long-term outcome should be considered when selecting treatment strategy.

Therefore, we conducted a multi-center registry survey in mainland China to explore the prevalence and characteristics of WO among outpatients with PD receiving L-dopa therapy. In addition, the factors considered by movement disorders specialists when making therapeutic choices for WO patients and the prevalence rate of dyskinesia were also investigated.

Subjects and methods

Survey setting

From May, 2012 to October, 2012, this cross-sectional, non-interventional registry study was performed simultaneously at 28 movement disorders clinics in tertiary hospitals located in capital cities with over 1,000 beds and the highest medical and academic levels in China. Patients with PD who were willing to participate, provided written informed consent, and met the entry criteria were enrolled at each clinic. During each visit, firstly, preliminary screening for eligible PD patients was done by study nurses before evaluated by the neurologists; Secondly, WOQ-9 and basic information registry form were completed by the informed patients, it could be completed with the aid of study nurse; Finally, comprehensive case report form (CRF) was recorded for patients with at least one positive item ($WOQ-9 \geq 1$). The final diagnoses of WO and dyskinesia were made by movement disorders specialists. WO was accepted as the shortening of the duration of L-dopa benefit to less than 4 h for every L-dopa dose given as reference [11]. Dyskinesia was defined as an involuntary choreatic movement involving the muscles of limbs, neck, trunk, or rarely face during the "on" period of patients who displayed prominent improvement of their parkinsonian symptoms [11].

The study was sponsored by Chinese Parkinson's Disease & Movement Disorders Society, Neurology Branch

of Chinese Medical Association, with approval from the Research Ethics Committee, Rui Jin Hospital affiliated to Shanghai Jiao Tong University School of Medicine, Shanghai, China.

Survey design

Subjects with clinical diagnosed PD based on the United Kingdom PD Brain Bank Criteria [12], were eligible for inclusion into the study if they had L-dopa medication at least 30 days and written informed consent. Patients with secondary parkinsonism and atypical parkinsonism syndrome (such as multiple system atrophy, progressive supranuclear palsy, dementia of lewy body, cortical basal ganglia degeneration, etc.) were excluded. Subjects were also excluded if they withdraw the survey underway.

9-item wearing-off questionnaire

For those eligible PD patients, WOQ-9 was completed, which covered both motor and non-motor symptoms of WO [13]. For each item, patients reported whether a symptom was present and whether it improved after the subsequent dose of anti-PD medications. If both were positive, one score was obtained. WOQ-9 defined the possible presence of WO as at least one score obtained.

Basic information registry form

For enrolled PD patients, demographic parameters such as age, gender, smoking, education, height, body weight and concomitant diseases were recorded on basic information registry form. Clinical characteristics collected included onset age, disease duration, L-dopa duration and medications history. L-dopa dosage and L-dopa equivalent dosage (LED) were calculated as reported by a previous report [14].

Case report form (CRF)

CRF was completed for PD patients with more than one positive item of WOQ-9. Besides information collected in basic registry form, the motor subscale of the Unified Parkinson's Disease Rating Scale (UPDRS-III) was used to evaluate motor function [15]. For patients with motor fluctuation, they were assessed during the "on" stage. Modified Abnormal Involuntary Movement Scale (mAIMS) was used to screen dyskinesia [16]. An Activities of Daily Living (ADL) questionnaire was completed by the patients or their caregivers or spouse, which included six physical and eight instrumental ADLs [17]. If movement disorders specialists considered the current treatment protocol of patients with WO needed adjustment, the factors concerned should be selected below: 1) long-term disease control and drug selection, 2) limited economic condition of patients, 3) better improvement of motor symptoms, 4) prior consideration of tolerance and safety of medications, 5) whether the drug was available in hospitals or

not, 6) the price of medications, 7) the patients' choices, 8) others. All the assessments were performed by movement disorders specialists during face-to-face interviews with the patients. Researchers were systematically trained before the investigation.

Statistical analysis

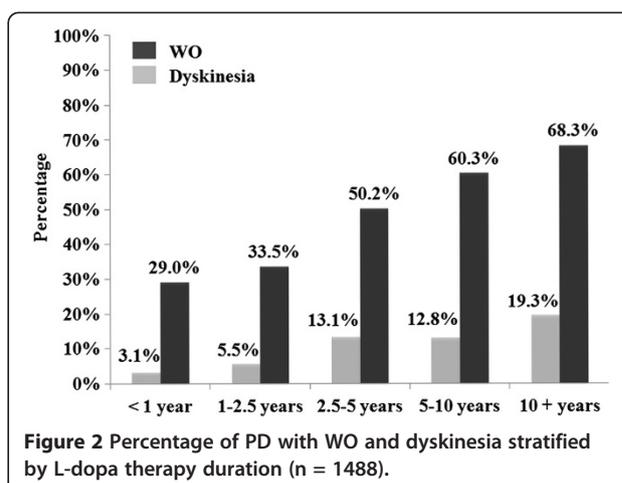
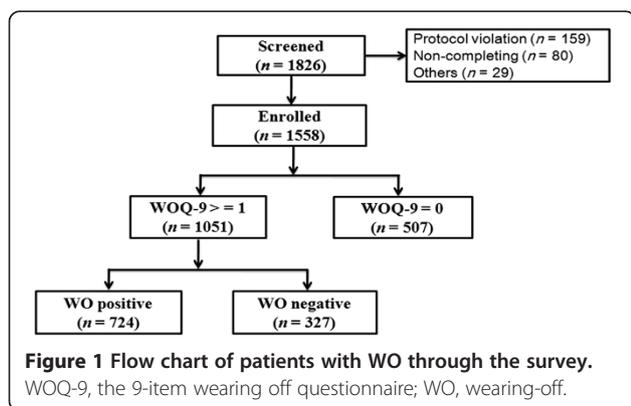
Statistical analysis was performed with SPSS, the overall and L-dopa duration specific prevalence rates of WO and dyskinesia were estimated and the corresponding 95% confidence interval (CI) were calculated based on the Poisson distribution. The independent sample *t* test and Chi-square analysis were employed for comparing group means and categorical data, respectively. Pearson correlation analysis was used for evaluating bivariate correlation. All *p* values were 2-sided and values < 0.05 were considered statistically significant.

Results

WO prevalence, characteristics and associated factors

Figure 1 described the screening and enrollment protocol through the survey. Of the 1,826 patients screened, 1,558 were enrolled in this study for further evaluation. A total of 268 were excluded, the majority of them were due to protocol violations (*n* = 159) and withdraw under way (*n* = 80). The protocol violation was caused by misdiagnosis of PD. Among these enrolled patients, a total of 507 had negative finding of WOQ-9. Thus, 1,051 with score ≥ 1 of WOQ-9 went into the evaluation stage of movement disorders specialists. The demographic and clinical data of enrolled patients and those with CRF evaluation were summarized in Additional file 1: Table S1. Finally, 724 patients were diagnosed with WO. The overall prevalence rate of WO was 46.5% (95% CI: 44.0% - 48.9%).

The prevalence of WO increased with the prolonged duration of L-dopa treatment (*p* = 0.0013), as shown in Figure 2. 29.0% PD patents with L-dopa duration less than one year experienced WO, the corresponding values were 33.5%, 50.2%, 60.3% and 68.3% for those with L-dopa duration 1–2.5 years, 2.5 - 5 years, 5–10 years and more than



10 years, respectively. The probability of detecting WO by movement disorders specialists was strongly associated with the positive response number of WOQ-9 (*p* = 0.0002).

Among these 724 confirmed patients with WO, the mean score of WOQ-9 was 3.8 (SD = 1.8). Overall, motor WO symptoms outweighed the non-motor WO symptoms (Figure 3). As the most common motor symptoms WO, slowness of movement was identified by up to 83.0% of patients. Whereas, the most common non-motor WO symptoms was pain/aching occupying 31.1% of subjects.

Compared with those without WO, patients with WO had more proportion of female patients (*p* < 0.0422), younger age (*p* < 0.0001), lower weight (*p* < 0.0090), earlier onset age (*p* < 0.0001), longer disease duration (*p* < 0.0001), longer L-dopa therapy duration (*p* < 0.0001), larger L-dopa dosage and LED (*p* < 0.0001), more severe UPDRS-III (*p* < 0.0001) and disabled ADL (*p* < 0.0001). Subgroup analysis for patients with CRF evaluation showed that 19.5% of patients with WO had dyskinesia as confirmed by movement disorders specialists, the proportion of which was significantly higher than that without WO (6.48%) (*p* < 0.0001). With respect to medications, there were more proportions of patients with WO receiving levodopa-benserazide (*p* < 0.0001), levodopa-carbidopa (*p* = 0.0002), Piribedil (*p* = 0.0013) and entacapone (*p* < 0.0001), relative to the WO negative group (Table 1).

Factors considered when adjusting therapeutic strategies for WO patients

Among 724 patients with WO, 55.7% (*n* = 403) of them needed therapy adjustment according to the opinion of movement disorders specialists. The factors concerned by movement disorder specialists for WO patients, in rank order, were as follows: better improvement of motor symptoms (*n* = 354, 87.8%), long-term disease control and drug selection (*n* = 288, 71.5%), prior consideration of tolerance and safety of medications (*n* = 54, 13.4%), limited

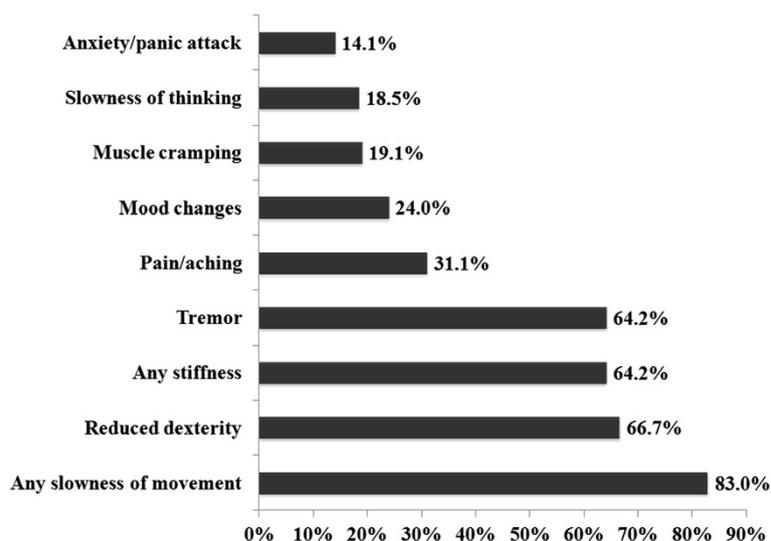


Figure 3 Frequency of motor and non-motor WO symptoms with respect to WOQ-9 among WO patients ($n = 724$).

economic condition of patients ($n = 35$, 8.7%), the patients' choices ($n = 14$, 3.5%) and the price of medications ($n = 11$, 2.7%).

Dyskinesia prevalence and associated factors

Among 1,558 enrolled patients, 160 of them were finally diagnosed with dyskinesia by movement disorders specialists. The overall prevalence rate of dyskinesia was 10.3% (95% CI: 8.8% - 11.8%). Dyskinesia prevalence rate was positively associated with L-dopa treatment duration ($p = 0.0072$) as shown in Figure 2.

Compared with those without dyskinesia, patients with dyskinesia had younger age ($p = 0.0016$), earlier onset age ($p < 0.0001$), lower weight ($p < 0.0001$), longer disease duration ($p < 0.0001$), longer L-dopa therapy duration ($p < 0.0001$), larger amount of L-dopa dosage and LED ($p < 0.0001$) and more disabled ADL ($p < 0.0001$). Subgroup analysis for patients with CRF evaluation showed that up to 88.5% of patients with dyskinesia had WO, which was significantly higher than those without dyskinesia (65.9%) ($p < 0.0001$). In terms of medications, there were more proportions of patients with dyskinesia receiving levodopa-carbidopa ($p < 0.0001$), piribedil ($p < 0.0001$) and entacapone ($p = 0.0003$), in comparison with those without dyskinesia. However, regarding the proportion of levodopa-benserazide, there was no difference between the two groups ($p = 0.9094$) (Table 1).

Discussion

This cross-sectional, multi-center survey showed that, for Chinese PD outpatients with L-dopa therapy, totally 46.5% and 10.3% suffered from WO and dyskinesia, respectively. Compared with those reported in other populations, the prevalence rate of WO was similar while the

prevalence rate of dyskinesia was obviously lower (Table 2). The underlying reasons may be as follows: 1) Methodological differences: in theory, results from current multi-center study and the other three studies [18-20] were more accurate than those from single center reports [2,4,5,11,21,22]. The current study revealed WO prevalence (46.5%) in China was a little bit higher than that in USA & Europe (motor fluctuation 43.9%) [19] and lower than that in Czech Republic(66.7%) [18]. 2) Variations on baseline characteristics of enrolled patients: Several prospective studies indicated L-dopa dosage and disease duration were two important predictive factors of the development of dyskinesia [3,23,24]. Both items of enrolled patients in the present study were lower than previous reports in North America, Europe and some Asian Countries, which may explain the low prevalence of dyskinesia in China. 3) Race factor and genetic predisposition: Olanow *et al.* found both WO and dyskinesia occurred more frequently in North American versus European sites [24]. It was reported in recent years that several genetic variations (such as COMT Val158Met [25], BDNF val66met [26], mu opioid receptor polymorphism and dopamine D2 receptor intronic dinucleotide repeat polymorphism [27]) were associated with the occurrence of motor complications, especially for dyskinesia. So the specific susceptible loci associated with motor complications in Chinese patients warrant further investigation.

Since WO is generally the first motor complication to develop, early identification of such condition is of great importance for the optimized therapy. This survey showed that approximately 67.5% of Chinese PD patients with more than one positive response of WOQ-9 were finally diagnosed with WO by movement disorders specialists, confirming that WOQ-9 was fit for clinical screening for

Table 1 Differences between enrolled patients with and without WO and dyskinesia

Items	WO			Dyskinesia		
	With (n = 724)	Without (n = 834)	P value	With (n = 160)	Without (n = 1398)	P value
Age, years			<.0001			0.0016
Mean ± SD	63.8 ± 9.6	66.5 ± 9.3		63.0 ± 9.5	65.5 ± 9.5	
n (missing)	710(14)	815(19)		156(4)	1369(29)	
Gender, Female			0.0422			0.1959
n (%)	347(47.9)	356(42.8)		80(50.0)	623(44.6)	
n (missing)	724(0)	832(2)		160(0)	1396(2)	
Height, cm			0.2230			0.2203
Mean ± SD	164.1 ± 7.8	164.6 ± 7.7		163.6 ± 8.3	164.4 ± 7.7	
n (missing)	717(7)	826(8)		157(3)	1386(12)	
Weight, kg			0.0090			<.0001
Mean ± SD	60.9 ± 11.1	62.3 ± 10.7		58.46 ± 10.12	62.01 ± 10.91	
n (missing)	720(4)	826(8)		158(2)	1388(10)	
Onset age, years			<.0001			<.0001
Mean ± SD	57.3 ± 9.9	62.3 ± 9.5		54.9 ± 9.9	60.5 ± 9.9	
n (missing)	711(13)	783(51)		156(4)	1338(60)	
Disease duration, years			<.0001			<.0001
Mean ± SD	6.7 ± 4.4	4.2 ± 3.7		8.2 ± 5.5	5.1 ± 4.0	
n (missing)	713(11)	796(38)		155(5)	1354(44)	
UPDRS-III*			<.0001			0.1678
Mean ± SD	19.0 ± 8.4	15.6 ± 6.7		18.8 ± 9.1	17.8 ± 7.9	
n (missing)	724(0)	327(0)		160(0)	891(0)	
L-dopa duration, years			<.0001			<.0001
Mean ± SD	5.1 ± 3.9	3.3 ± 3.5		5.9 ± 4.7	3.9 ± 3.7	
n (missing)	703(21)	785(49)		151(9)	1337(61)	
L-dopa dosage			<.0001			<.0001
Mean ± SD	478.5 ± 247.0	355.4 ± 191.5		581.7 ± 311.4	393.8 ± 207.7	
n (missing)	724(0)	834(0)		160(0)	1398(0)	
≤400 mg/d n (%)	359(37.2)	607(62.8)	<.0001	52(5.4)	914(94.6)	<.0001
400-600 mg/d n (%)	234(56.5)	180(43.5)		54(13.0)	360(87.0)	
>600 mg/d n (%)	131(73.6)	47(26.4)		54(30.3)	124(69.7)	
LED			<.0001			<.0001
Mean ± SD	558.4 ± 281.7	414.4 ± 216.0		707.8 ± 336.9	455.4 ± 234.9	
n (missing)	724(0)	834(0)		160(0)	1398(0)	
ADL*			<.0001			<.0001
Mean ± SD	28.4 ± 12.9	21.0 ± 9.6		31.3 ± 13.6	25.1 ± 12.0	
n (missing)	721(3)	326(1)		160(0)	887(4)	
Medications for PD, n (%)						
Levodopa-benserazide	647(89.4)	682(81.8)	<.0001	136(85.0)	1193(85.3)	0.9094
Levodopa-carbidopa	285(39.4)	253(30.3)	0.0002	87(54.4)	451(32.3)	<.0001
Pramipexole	245(33.8)	266(31.9)	0.4147	58(36.3)	453(32.4)	0.3262
Piribedil	196(27.1)	168(20.1)	0.0013	66(41.3)	298(21.3)	<.0001

Table 1 Differences between enrolled patients with and without WO and dyskinesia (Continued)

Entacapone	152(21.0)	109(13.1)	<.0001	43(26.9)	218(15.6)	0.0003
Selegiline	17(2.4)	32(3.8)	0.0931	4(2.5)	45(3.2)	0.6216
Trihexyphenidyl	101(13.9)	114(13.7)	0.8725	17(10.6)	198(14.2)	0.2190
Amantadine	174(24.0)	182(21.8)	0.3000	46(28.8)	310(22.2)	0.0606

*based on patients with complete CRF; WO Wearing-off, UPDRS Unified Parkinson's Disease Rating Scale, LED, Levodopa equivalent dosage, ADL Activities of Daily Living.

WO, as recommended by MDS in 2011 [8]. With respect to WO phenomenology, totally, motor WO symptoms outweighed the non-motor WO symptoms, as shown in the present survey and Western Countries' surveys. Movement slowness was the most common motor WO symptoms, whereas, pain/aching was the most common non-motor WO symptoms in our results. This may be a little bit difference from those in Western Countries. Stacy and co-workers in 2005 found tremor and tiredness were the most frequent motor and non-motor fluctuation symptom, respectively [19]. The potential reason of this discrepancy needs to be further investigated. However, both of these symptoms may provide important information for the identification of WO in clinical practice. It was reported that non-motor fluctuation may involve a greater degree of disability than motor fluctuation [28]. Therefore, more attention should be paid for the recognition of non-motor fluctuation in future research.

Our survey indicated that young onset age, low body weight, large amount of LED, long duration of disease and L-dopa therapy were associated with motor complications in Chinese PD patients with impaired ADL. Other than those with dyskinesia, there were more female and severe motor disabled patients with WO. All these results were consistent with previous cross-sectional investigations by others and a recent prospective study reported by Olanow and co-workers [24]. Based on these studies, clinicians should initiate L-dopa treatment for PD patients

with low doses and increase the dosage in small increments in the long run. Besides, it may also be important to consider the weight of patients when prescribing L-dopa. In terms of medications, patients with motor complications took more proportions of L-dopa, piribedil and entacapone. Since this was a cross-sectional study, a proportion of patients may have already employed the adjunctive therapy (such as piribedil or entacapone) for their existed motor complications, we could not draw a conclusion whether these alternative drugs were risk or protective factors of motor complications in the present survey.

As expected, better improvement of motor symptoms (87.8%) and long-term disease control and drug selection (71.5%) were the two frequent considered factors when movement disorders specialists adjusted therapeutic strategies for patients with WO. This may be partially benefited from the clinical practice of Chinese PD treatment guidelines, since published in 2006 and 2009, respectively [9]. The L-dopa treatment concept—using the lowest dose that provides the satisfactory clinical control to delay the occurrence of motor complication, especially among PD patients with young onset age, influenced more and more clinical doctors with its popularity.

There were some limitations of our study. Firstly, patients with no positive response of WOQ-9 did not go to neurologists' evaluation in the present survey. Besides, only choreatic dyskinesias were counted which could leave out other types of dyskinesia such as dystonic

Table 2 Serial epidemiological studies of PD patients with WO and dyskinesia in Asian and Caucasians

Countries	PD sample	Design	WO prevalence (%)	Dyskinesia prevalence (%)	References
UK	87	Community based	40.0(MF)	28	Schrag A, et al. <i>Brain</i> , 2000 [2]
China	63	Hospital based, single center	44.4	17.5	Liu CF, et al. <i>Chin J Neurol</i> , 2003 [5]
USA & Europe	289	Hospital based, multi-center	43.9*(MF)	-	Stacy M, et al. <i>Mov Disord</i> , 2005 [19]
Turkey	555	Hospital based, single center	46.3	30.1	Benbir G, et al. <i>Clin Neurol Neurosurg</i> , 2006 [11]
Chile	124	Hospital based, single center	52.0	47.2	Juri-Claveria C, et al. <i>Rev Neurol</i> , 2007 [21]
Hong Kong	98	Hospital based, single center	74.5(MF)	77.6	Kum WF, et al. <i>J Clin Neurosci</i> , 2009 [4]
Czech Republic	563	Hospital based, multi-center	66.7	-	Bares M, et al. <i>J Neural Transm</i> , 2012 [18]
Japan	1453	Hospital based, single center	44.7	-	Yoritaka A, et al. <i>Parkinsonism Relat Disord</i> , 2013 [22]
Italy	617	Hospital based, multi-center	56.9	-	Stocchi F, et al. <i>Parkinsonism Relat Disord</i> , 2014 [20]
China	1558	Hospital based, multi-center	46.5	10.3	Present study

[†]WO diagnosis according to UPDRS-IV, MF Motor fluctuation.

dyskinesias. Therefore, the true prevalence rate of WO and dyskinesia might be a little bit underestimated. Secondly, as a cross-sectional study, only associated factors of motor complications could be found in the present investigation. Since only patients with more than one point of WOQ-9 had UPDRS motor evaluation, we did not perform multiple logistic regression analysis to explore the independent influential factor of WO and dyskinesia. To better clarify its independent predictive factors, prospective cohort studies are needed in future.

Conclusions

We performed the first multi-center registry survey of WO among Chinese patients with PD on L-dopa therapy. The prevalence rate of WO found in our study was similar to, while dyskinesia was relatively lower than those in other Asian and Caucasian Countries, and different reasons such as methodological difference, L-dopa dosage and genetic susceptibility may contribute to this result.

Appendix

The following principle investigators participated in the study (Sequence arrangement according to a descending order of included case number): Sheng-Di Chen, Rui Jin Hospital affiliated to Shanghai Jiao Tong University School of Medicine; Ming Shao, The First Affiliated Hospital of Guangzhou Medical College; Tao Feng, Beijing Tiantan Hospital affiliated to Capital Medical University; Wei-Guo Liu, Nanjing Brain Hospital affiliated to Nanjing Medical University; Xiao-Guang Luo, The First Hospital of China Medical University; Xiao-Chun Chen, Union Xiehe Hospital, Fujian Medical University; An-Mu Xie, The Affiliated Hospital of Medical College Qingdao University; Chun-Feng Liu, The Second Affiliated Hospital of Soochow University; Zhen-Guo Liu, Xinhua Hospital affiliated to Shanghai Jiao Tong University School of Medicine; Yi-Ming Liu, Qilu Hospital of Shandong University; Jian Wang, Huashan Hospital affiliated to Fudan University; Hui-Fang Shuang, West China Hospital, Sichuan University; Biao Chen, Xuanwu Hospital of Capital Medical University; Bei-Sha Tang, Xiangya Hospital, Central South University; Xing-Yue Hu, Shaoyifu Hospital affiliated to Zhejiang University School of Medicine; Li-Juan Wang, Guangdong General Hospital; Bao-Rong Zhang, The Second Affiliated Hospital of Zhejiang University School of Medicine; Min Ye, Nanjing BenQ medical Center; Hai-Bo Chen, Peking Hospital; Xin-Hua Wan, Peking Union Medical College Hospital; Ming-Wei Wang, The First Hospital of Hebei Medical University; Xian-Wen Chen, The First Affiliated Hospital of Anhui Medical University; Yan Chen, General Hospital Affiliated to Tianjing medical University; Guo-Guang Peng, The First Affiliated Hospital of Chongqing Medical University; Zhen-Fu Wang,

The General Hospital of Chinese People's Liberation Army; Ping-Yi Xu, The First Affiliated Hospital, Sun Yat-Sen University; Sheng-Gang Sun, Union Hospital of Tongji Medical College of Huazhong University of Science and Technology; Xiang-Ru Sun, The First Hospital affiliated to Beijing University.

Additional file

Additional file 1: Table S1. Demographic and basic clinical information of enrolled patients and those with complete CRF.

Abbreviations

PD: Parkinson's disease; WO: Wearing off; WOQ-9: 9-item wearing off questionnaire; COMT: Catechol-O-methyltransferase; MAO-B: Monoamine oxidase-B.

Competing interests

The authors declare that they have no competing interests.

Authors' contributions

Study concept and design: S-D C. Acquisition of data: WC, QX, MS, TF, W-G L, X-G L, X-Chun C, A-M X, C-F L, Z-G L, Y-M L and JW. Statistical analysis: H&J CRO International, Inc. Analysis and interpretation of data: WC, QX and S-D C. Drafting of the manuscript: WC and QX. Critical revision of the manuscript for important intellectual contents: S-D C. Study supervision: QX, S-D C. All authors read and approved the final manuscript.

Acknowledgements

The authors would like to acknowledge and thank all the patients, care providers, and investigators who participated in this survey for their commitment to the project. We also appreciated the statistical analysis from H&J CRO International, Inc. This study was supported by Novartis China. This work was also supported by grants from the National Program of Basic Research (2011CB504104) of China and National "Twelfth Five-Year" Plan for Science & Technology Support (2012BA110B03).

Author details

¹Department of Neurology, Rui Jin Hospital affiliated to Shanghai Jiao Tong University School of Medicine, Rui Jin 2nd Road 197, Shanghai 200025, China. ²Department of Neurology, The First Affiliated Hospital of Guangzhou Medical College, Guangzhou, China. ³Department of Neurology, Beijing Tiantan Hospital affiliated to Capital Medical University, Beijing, China. ⁴Department of Neurology, Nanjing Brain Hospital affiliated to Nanjing Medical University, Nanjing, China. ⁵Department of Neurology, The First Hospital of China Medical University, Shenyang, China. ⁶Department of Neurology, Union XieHe Hospital, Fujian Medical University, Fuzhou, China. ⁷Department of Neurology, The Affiliated Hospital of Medical College Qingdao University, Qingdao, China. ⁸Department of Neurology, The Second Affiliated Hospital of Soochow University, Suzhou, China. ⁹Department of Neurology, Xinhua Hospital affiliated to Shanghai Jiao Tong University School of Medicine, Shanghai, China. ¹⁰Department of Neurology, Qilu Hospital of Shandong University, Jinan, China. ¹¹Department of Neurology, Huashan Hospital affiliated to Fudan University, Shanghai, China.

Received: 13 October 2014 Accepted: 26 November 2014

Published: 5 December 2014

References

1. Stacy M: The wearing-off phenomenon and the use of questionnaires to facilitate its recognition in Parkinson's disease. *J Neural Transm* 2010, **117**:837–846.
2. Schrag A, Quinn N: Dyskinesias and motor fluctuations in Parkinson's disease. A community-based study. *Brain* 2000, **123**:2297–2305.
3. Ahlskog JE, Muenter MD: Frequency of levodopa-related dyskinesias and motor fluctuations as estimated from the cumulative literature. *Mov Disord* 2001, **16**:448–458.

4. Kum WF, Gao J, Durairajan SS, Man SC, Xie LX, Lu JH, Fong WL, Li M: **Risk factors in development of motor complications in Chinese patients with idiopathic Parkinson's disease.** *J Clin Neurosci* 2009, **16**:1034–1037.
5. Liu CF, Yin WH, Luo WF: **Factors of dyskinesias and motor fluctuations in Parkinson's disease.** *Chin J Neurol* 2003, **36**:411–413.
6. Stacy MA, Murphy JM, Greeley DR, Stewart RM, Murck H, Meng X: **The sensitivity and specificity of the 9-item Wearing-off Questionnaire.** *Parkinsonism Relat Disord* 2008, **14**:205–212.
7. Chan A, Cheung YF, Yeung MA, Yeung J, Chung TH, Tsang KL, Chan J, Lau C, Kwan P, Kuo SH, Mok V: **A validation study of the Chinese wearing off questionnaire 9-symptom for Parkinson's disease.** *Clin Neurol Neurosurg* 2011, **113**:538–540.
8. Antonini A, Martinez-Martin P, Chaudhuri RK, Merello M, Hauser R, Katzenschlager R, Odin P, Stacy M, Stocchi F, Poewe W, Rascol O, Sampaio C, Schrag A, Stebbins GT, Goetz CG: **Wearing-off scales in Parkinson's disease: critique and recommendations.** *Mov Disord* 2011, **26**:2169–2175.
9. Chen W, Chen S, Xiao Q, Wang G, Chen SD: **Current clinical practice for Parkinson's disease among Chinese physicians, general neurologists and movement disorders specialists: a national survey.** *BMC Neurol* 2012, **12**:155.
10. Muller T: **Drug therapy in patients with Parkinson's disease.** *Transl Neurodegener* 2012, **1**:10.
11. Benbir G, Ozekmekci S, Apaydin H, Delil S, Erginoz E: **A hospital-based study: risk factors in development of motor complications in 555 Parkinson's patients on levodopa therapy.** *Clin Neurol Neurosurg* 2006, **108**:726–732.
12. Hughes AJ, Daniel SE, Kilford L, Lees AJ: **Accuracy of clinical diagnosis of idiopathic Parkinson's disease: a clinico-pathological study of 100 cases.** *J Neurol Neurosurg Psychiatry* 1992, **55**:181–184.
13. Stacy M, Hauser R, Oertel W, Schapira A, Sethi K, Stocchi F, Tolosa E: **End-of-dose wearing off in Parkinson disease: a 9-question survey assessment.** *Clin Neuropharmacol* 2006, **29**:312–321.
14. Tomlinson CL, Stowe R, Patel S, Rick C, Gray R, Clarke CE: **Systematic review of levodopa dose equivalency reporting in Parkinson's disease.** *Mov Disord* 2010, **25**:2649–2653.
15. Richards M, Marder K, Cote L, Mayeux R: **Interrater reliability of the Unified Parkinson's Disease Rating Scale motor examination.** *Mov Disord* 1994, **9**:89–91.
16. Guy W, AIMS: *ECDEU Assessment Manual for Psychopharmacology.* Washington, DC: Government Printing Office; 1976:534.
17. Lawton MP, Brody EM: **Assessment of older people: self-maintaining and instrumental activities of daily living.** *Gerontologist* 1969, **9**:179–186.
18. Bares M, Rektorova I, Jech R, Farniková K, Roth J, Růžička E, Kaňovský P, Rektor I, Pavlík T, Uhlířová L, Vydlák J: **Does WOOQ-9 help to recognize symptoms of non-motor wearing-off in Parkinson's disease.** *J Neural Transm* 2012, **119**:373–380.
19. Stacy M, Bowron A, Guttman M, Hauser R, Hughes K, Larsen JP, LeWitt P, Oertel W, Quinn N, Sethi K, Stocchi F: **Identification of motor and nonmotor wearing-off in Parkinson's disease: comparison of a patient questionnaire versus a clinician assessment.** *Mov Disord* 2005, **20**:726–733.
20. Stocchi F, Antonini A, Barone P, Tinazzi M, Zappia M, Onofri M, Ruggieri S, Morgante L, Bonuccelli U, Lopiano L, Pramstaller P, Albanese A, Attar M, Posocco V, Colombo D, Abbruzzese G: **Early DEtection of wEaring off in Parkinson disease: The DEEP study.** *Parkinsonism Relat Disord* 2014, **20**:204–211.
21. Juri-Claveria C, Aguirre-M C, Viviani-G P, Chana-Cuevas P: **Risk factors associated with the development of motor complications in Parkinson's disease. A study in a Chilean population.** *Rev Neurol* 2007, **45**:77–80.
22. Yoritaka A, Shimo Y, Takanashi M, Fukae J, Hatano T, Nakahara T, Miyamoto N, Urabe T, Mori H, Hattori N: **Motor and non-motor symptoms of 1453 patients with Parkinson's disease: prevalence and risks.** *Parkinsonism Relat Disord* 2013, **19**:725–731.
23. Hauser RA, McDermott MP, Messing S: **Factors associated with the development of motor fluctuations and dyskinesias in Parkinson disease.** *Arch Neurol* 2006, **63**:1756–1760.
24. Warren OC, Kieburz K, Rascol O, Poewe W, Schapira AH, Emre M, Nissinen H, Leinonen M, Stocchi F: **Factors predictive of the development of Levodopa-induced dyskinesia and wearing-off in Parkinson's disease.** *Mov Disord* 2013, **28**:1064–1071.
25. de Lau LM, Verbaan D, Marinus J, Heutink P, van Hilten JJ: **Catechol-O-methyltransferase Val158Met and the risk of dyskinesias in Parkinson's disease.** *Mov Disord* 2012, **27**:132–135.
26. Foltynie T, Cheeran B, Williams-Gray CH, Edwards MJ, Schneider SA, Weinberger D, Rothwell JC, Barker RA, Bhatia KP: **BDNF val66met influences time to onset of levodopa induced dyskinesia in Parkinson's disease.** *J Neurol Neurosurg Psychiatry* 2009, **80**:141–144.
27. Strong JA, Dalvi A, Revilla FJ, Sahay A, Samaha FJ, Welge JA, Gong J, Gartner M, Yue X, Yu L: **Genotype and smoking history affect risk of levodopa-induced dyskinesias in Parkinson's disease.** *Mov Disord* 2006, **21**:654–659.
28. Witjas T, Kaphan E, Azulay JP, Blin O, Ceccaldi M, Pouget J, Poncet M, Chérif AA: **Nonmotor fluctuations in Parkinson's disease: frequent and disabling.** *Neurology* 2002, **59**:408–413.

doi:10.1186/2047-9158-3-26

Cite this article as: Chen et al.: Prevalence of wearing-off and dyskinesia among the patients with Parkinson's disease on levodopa therapy: a multi-center registry survey in mainland China. *Translational Neurodegeneration* 2014 **3**:26.

Submit your next manuscript to BioMed Central and take full advantage of:

- Convenient online submission
- Thorough peer review
- No space constraints or color figure charges
- Immediate publication on acceptance
- Inclusion in PubMed, CAS, Scopus and Google Scholar
- Research which is freely available for redistribution

Submit your manuscript at
www.biomedcentral.com/submit

