Introduction

Pisa syndrome (PS) is a postural deformity which clinically presents as a marked lateral flexion of the trunk with a bending degree greater than 10° or 15°, typically mobile in nature and resolves at supine position. It is characterized by dystonia, and abnormal and sustained involuntary muscle contraction which may cause twisting or jerking movements of the body or a body part [1-3,17] (Figure 1). Ekbom, et al. originally described this entity which was then associated with butyrophenones [4]. It has been described in Parkinson's disease (PD) either related to disease itself or secondary to levodopa or COMT inhibitors [9], atypical parkinsonism [5] and other neurodegenerative disorders like AD [6], DLB [7], HD [8], SSPE [10]. PS is linked to almost all dopaminergic agents, antipsychotics, valproate, benzodiazepines, lithium [11]. PS has been described in normotensive hydrocephalus [12,13], subdural hematoma [14], late complication of pallidotomy in a patient with PD [15] or even idiopathic [16].

Epidemiology

Prevalence of PS is not well known because of the lack of clear diagnostic criteria. However, studies done previously reported that it may range from 2% to 90%. Tinazzi, et al. reported prevalence of 8.8% in PS patients with PD. The authors also reported that PS patients in their study were older, had lower body mass index, longer duration of disease in years, higher stages of disease, poorer quality of daily life, more frequent falls as well as occurrence of “veering gait” (the progressive deviation toward one side when patient walked forward and backward with eyes closed). The most frequently associated medical conditions were osteoporosis and arthrosis [3].

Prevalence rate of 8.3% (9.3% in women, 6.4% in men)
was found in a Canadian study in which 133 patients were followed up for 2 days-3 months after typical neuroleptic exposure [18], contrary to a German study with psychiatric patients receiving antipsychotics where prevalence was reported 0.037% [19]. Risk factors associated in this study were females, old age, neuroleptic exposure and organic brain disorder. In a surveillance study which looked into the association of cholinesterase inhibitors and PS, mean (SD) age of 51 cases was 74.6 (8.2) and female: male ratio of 2:1 was reported [20]. In a single-center study from Italy, 26 patients were reported with lateral trunk deviation in the cohort of 1,400 patients with parkinsonism [21]. Scoliosis is associated with PD and parkinsonism and can also be confused with PS [22-24].

**Pathophysiology**

The pathophysiology of PS is not clear. Accepted hypothesis till date are divided into central and peripheral. Central mechanisms mostly involve alteration in sensory-motor integration pathways, basal ganglia dysfunction secondary to cholinergic-dopaminergic imbalance whereas peripheral hypothesis is associated with anatomical changes in the musculoskeletal system [1,17,29,31]. Cognitive processes and dysfunction also might be related to PS [26]. Altered visual-spatial functions and vertical perception deficits might represent a typical feature of PD patients with PS [26,27]. PS in PD patients may involve the nondopaminergic pathways as evident therapy-resistant symptom exists in this subset population [30].

Studies from animal models till date only suggests that there is a role of an asymmetric functioning of the nigrostriatal dopaminergic projections in PS [28]. One study reported that, levodopa aggravated lateral flexion of neck and trunk post unilateral pallidotomy which may be a delayed phenomenon [32]. Recently, role of vestibular function in maintaining posture was studied in parkinsonian patients. They found out unilateral peripheral vestibular hypofunction was present in all patients with lateral trunk flexion and the vestibular hypofunction was ipsilateral to the leaning side and contralateral to the most affected parkinsonian side in all the patients [33].

A dystonic activity might play a vital role in determining the bending ipsilaterally to PS and the contralateral excessive muscle activation which may be a compensatory mechanism [34]. An abnormal tonic hyperactivity on the side of the trunk’s deviation in abdominal oblique muscles was reported in PD patients with PS [35]. Frazzitta, et al. reported asymmetric ability to generate maximal voluntary force of the external oblique muscles supporting a central desynchronization of axial muscles as a significant contributor for the bending of the spine in erect position [36]. Degenerative spinal disease, soft tissue and muscle changes should not be forgotten while managing patients in clinical practice. Doherty, et al. in their study reported abnormal posture can be combination of degeneration of muscles and soft tissue secondary to dystonia which may be present from before or possibly a complex impairment of proprioceptive motor control in PD patients with PS [37].

**Pathology**

Hozumi and colleagues reported an autopsy case of a 62-year-old Japanese woman diagnosed as MSA-P with PS. This patient had a clinical history of leaning towards her right side. Histopathological findings of brain showed neuronal loss and astrogliosis in the putamina more severe on the right side as compared to the left [38]. However a recent pathology report of a PD patient with PS found no significant basal ganglia asymmetry or brain stem involvement [39]. Peripheral pathology may be musculoskeletal in origin with myopathic alterations in paraspinal muscles [29].

**Diagnosis**

Diagnosis of PS requires at least 10° lateral flexion, which can be completely alleviated by passive mobilization or lying in a supine position [3,17]. Diagnosis of PS is based on the clinical evaluation of lateral displacement of the trunk, which is the first sign of postural misalignment usually reported by patients and their caregivers. Tracking can be done meticulously in day care clinical practice by goniometer, inclinometer or even by smartphone applications. X-rays in standing and supine positions is deemed necessary however standing is the most accurate method to assess the angle of curvature in the coronal and sagittal planes according to the Cobb angle to rule out other diagnoses [40,41] (Figures 2,3).
**Clinical manifestation**

PS can develop in acute, subacute or chronic fashion [42-44]. PS patients generally lean towards one side while sitting, standing, and walking and also can have impaired perception of their vertical position awareness [17]. Recent exposure to antipsychotics and other medications may be the part of acute and subacute forms which benefits the most from anticholinergic treatment, however those with chronic exposure whose onset of the syndrome is insidious are less responsive to anticholinergic treatment [44-46]. This led Prahraj, et al. to classify PS into acute and tardive type [47]. However it is also reported that patients with chronic advanced type have a tendency to tilt to one side when sitting in a chair with subsequent lateral flexion during walking [48]. When the deformity worsens over time it can lead to dyspnea, low back pain [49], or unsteadiness leading to falls [50,51].

**Differential diagnosis**

PS can be differentiated from tardive dystonia [44], which is triggered by intake of dopamine receptor blockers such as antipsychotics for other causes, typically after chronic treatment of around 5 years, which may be persistent or even permanent even when the offending drug is discontinued [52]. Scoliosis is defined as the lateral curvature of the spine with a Cobb angle of 10 degrees or more in the coronal plane as measured on a radiograph [53]. This impairment in scoliosis is monoplaner and resolves when the primary abnormality is treated [54]. Muscle inflammatory disease like focal myositis [55], neuromuscular junction disorders specifically myasthenia gravis [56], and facioscapulohumeral dystrophy [57] should also be kept in mind while considering the diagnosis of PS.

**Treatment**

**Pharmacological approaches**: Drug-induced PS predominantly develops in women and older patients with organic brain changes [11,66]. In patients who has developed PS acute or subacutely it is mandatory for clinicians and neurologists to find out if there is any recent changes in medication dose or any new medication added to either modify or remove it respectively because PS in non-PD patients has been frequently related to the use of dopamine receptor blockers or cholinesterase inhibitors [63-65].

Axial symptoms and PS respond poorly to dopaminergic therapy, however other dopaminergic agents like levodopa in high dosage can be tried in patients where PS appears as a motor complication during the off periods [58,61]. Other agents which can be used in PS may include anticholinergics [16] and novel antipsychotics without interference with dopaminergic receptors like clozapine [59], quetiapine, however clozapine has been reported in literature to cause PS [60].

Clinicians should be very precautious when prescribing atypical antipsychotics because they can induce and aggravate

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**Table 1**: Causes of PISA syndrome.

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<tr>
<th>Neurodegenerative diseases</th>
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<td>Parkinson’s disease</td>
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<td>Progressive supranuclear palsy</td>
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<td>Huntington’s disease</td>
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<td>Multiple system atrophy</td>
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**Drugs**

- Levodopa combinations (levodopa/carbidopa, levodopa/benserazide, levodopa/carbidopa/entacapone)
- Antipsychotics (typical and atypical)
- Lithium
- Cholinesterase inhibitors
- Ergot derivative—pergolide
- Nonergot derivatives—pramipexole and ropinirole
- Monoamine oxidase-B inhibitor—rasagiline
- Antiemetics
- Selective serotonin reuptake inhibitors
- Tricyclic antidepressants
- Benzodiazepines
- Valproate
- Idiopathic form

**Table 2**: Treatment modalities for PS.

- Pharmacological: High dose levodopa, anticholinergics, quetiapine, clozapine (cautiously), Istradefylline.
- Botulinum Toxin (BoNT)
- Surgical treatment: DBS, Spinal surgery
- Orthotics
- Rehabilitation

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PS [62]. Slower withdrawal of dopamine agonists might be another option as evidenced in patients with antecolls [17]. One recently published research reported that postural deformities caused by dopamine agonists generally improve less than two weeks after dopamine agonist withdrawal. Istradeffyline which is used as an add-on treatment to levodopa/carbidopa in adults with PD experiencing “off “ episodes can be a potential therapeutic option in postural deformities like PS [83].

**Botulinum toxin:** Excessive muscular hyperactivity in PS can be treated with botulinum toxin (BoNT) [67]. In a blinded crossover study by Bonanni, et al. six patients treated with BoNT showed an improvement between 50% and 87.5%. In one patient, only subjective benefit was reported, while two patients did not report any benefit [21]. Injection in paraspinal muscles should be avoided, and top most priority should be given to infiltrate external oblique muscle [36]. Iliopsoas and the rectus abdominis were the most frequently used muscle for injection in study conducted by Tassorelli, et al. [68]. Dupeyron, et al. reported complete and 1 year lasting resolution of PS after BoNT injection in the quadratus lumborum muscle [69]. Two unique studies also found out BoNT is useful to enhance the effect of rehabilitation treatment in PS patients [68,70].

**Orthotics:** Spinal short segment decompresion can be considered in appropriate candidates with spinal stenosis with radiculopathy or myelopathy as supported by a study on patients having PD with camptocormia [71].

**Rehabilitation:** Capecci, et al. conducted a single blind randomized clinical trial in which they applied postural rehabilitation for 4 weeks on 13 patients with PD and postural abnormalities including PS with 1-month follow-up. Six patients in the treatment group also had Kinesio taping strips applied to their trunk muscles. All treated patients showed a significant improvement in trunk posture in both the sagittal and coronal planes compared with baseline. Moreover, they showed an improvement in measures of gait and balance. They also concluded that combination of active posture correction and trunk movements, muscle stretching, and proprioceptive stimulation may usefully impact axial symptoms in PD. Repeated training was recommended to these patients to avoid waning of the effect. No differences were found between patients who received postural rehabilitation plus Kinesio taping bands compared with those receiving postural rehabilitation only [72].

One study also demonstrated significant improvements in axial posture and trunk mobility through the 4-week rehabilitation programme in patients with lateral trunk flexion in PD. However, the benefit was not sustained and diminished after a few months [73]. Frazzitta, et al. in their study also reported early rehabilitation before development of postural deformities in PD patients emphasizing specially on stretching exercises for the external oblique and paraspinal muscles [36]. Motor rehabilitation is an effective tool for PD patients with PS [68,70,74]. A structured exercise program to strengthen contralateral paraspinal muscles may be a promising strategy [34].

**Surgical treatment:** Deep brain stimulation (DBS) of subthalamic nucleus (STN) does not have a good outcome in amelioration of PD axial symptoms, freezing of gait, falls, including posture and postural instability [75]. Stefani, et al. studied stimulation of pedunculopontine nucleus (PPN) in severe PD. They suggested patients with advanced PD, PPN-DBS associated with standard STN-DBS may be useful in improving gait and in optimizing the dopamine-mediated ON-state, particularly in those whose response to STN only DBS has deteriorated over time [76].

Shih, et al. reported improvement of gait and lean in PS patient with contralateral pedunculopontine stimulation [77]. Ricciardi, et al. reported about a case where stimulator electrodes were implanted in PPN ipsilateral to the side of the bend because of its role in promoting atonia. However, improvement did not sustain over time [78]. Spinal surgery may be useful in complex spinal deformities where conservative management has failed however selection procedure should be individualized and meticulous [79].

**Discussion**

Postural deformities in PD is not uncommon in clinical practice but sometimes there can be an overlap of two conditions in the same patient like camptocormia and PS which makes situation for clinicians more difficult in terms of treatment. The pathophysiological mechanism may be same or at least closely related [44]. The etiology of this condition may be difficult to find as there is no consensus defenition of the abnormalities in the coronal plane.

PS can be acute, subacute or can develop slowly with the evolvement of underlying disease. Acute and subacute forms may be dystonia like phenomenon which needs more understanding at this point of time [81]. Chronic changes may be changes induced over long time affecting muscle and bones [44,50]. More recent reports suggests the association between PS and specific cognitive alterations, implying a potential contribution of cortical and subcortical dysfunctions in the pathophysiology of PS [82]. Gradual withdrawal of drugs causing PS can be beneficial along with use of anticholinergics in patients exposed to antipsychotics due to any reason [46,80]. Other options may be dopaminergic agents like levodopa, BoNT, orthotics, rehabilitation and surgery in the form of DBS or spinal surgery.

**Conclusion**

PS is considered to be an overlooked condition with limited epidemiologic data in clinical practice. It is common yet not specific to patients with PD only. It has a significant impact
on quality of life and daily functioning of an individual. A complex interplay between central basal ganglia dysfunction, together with proprioceptive disintegration and altered cognitive processing, is usually required for the development of PS. Physical symptoms like low back pain, dyspnea and fall may follow if not treated appropriately. Early recognition is must because it can be reversed at an acute or subacute stage. Future studies with an intention to gather enormous epidemiological data, understanding of pathophysiological mechanisms, consensus on diagnostic criteria followed by case to case based management is mandatory.

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**References**


